Hertzog

10/688,243

08/17/2004

STIC SEARCH

=> file reg

FILE 'REGISTRY' ENTERED AT 11:38:40 ON 17 AUG 2004 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2004 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 16 AUG 2004 HIGHEST RN 727651-15-2 DICTIONARY FILE UPDATES: 16 AUG 2004 HIGHEST RN 727651-15-2

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

=> d his

(FILE 'HOME' ENTERED AT 09:56:55 ON 17 AUG 2004)

FILE 'HCA' ENTERED AT 09:59:07 ON 17 AUG 2004 E US20040086443/PN

L1 1 S E3 SEL L1 RN

FILE 'REGISTRY' ENTERED AT 09:59:40 ON 17 AUG 2004

L2 10 S E1-E10

FILE 'LCA' ENTERED AT 09:59:48 ON 17 AUG 2004

L3 2119 S (RETARDER? OR INHIBITOR? OR SUPPRESSOR? OR SCAVENG?)

L4 1899 S SULFUR# OR SULPHUR# OR (SULFUR# OR SULPHUR#) (N) HYDRIDE## OR H

FILE 'HCA' ENTERED AT 10:04:28 ON 17 AUG 2004

L5 946220 S L3

L6 666696 S L4

L7 3153 S L5(3N)L6

FILE 'LCA' ENTERED AT 10:05:13 ON 17 AUG 2004

L8 65 S SULFHYDRYL# OR SULPHYDRIL#

L9 1597 S RECOVER? OR RECLAIM? OR RECLAMAT? OR RE(W) (COVER? OR CLAMAT?

L10 5286 S LIQUID# OR LIQ# OR MOLTEN# OR FLUID? OR MELTED? OR LIQUIF? OR

L11 3282 S SUSPEN? OR DISPERS? OR COLLOID? OR EMULS? OR MICROEMULS? OR S

FILE 'HCA' ENTERED AT 10:12:44 ON 17 AUG 2004

L12 22428 S L8

L13 20487 S L4(3N)(L10 OR L11) L14 85 S L7 AND L12

FILE 'HCA' ENTERED AT 10:14:14 ON 17 AUG 2004

FILE 'REGISTRY' ENTERED AT 10:14:35 ON 17 AUG 2004

- L15 6 S L2 AND 1-10/NR
- L16 4 S L2 NOT L15
- L17 2 S L16 AND SULFUR
- L18 1 S L17 AND 1-3/H
- L19 1 S L17 NOT L18
- L20 2 S L16 NOT L17
- L21 1 S L15 AND CYCLOHEXENE
- L22 7 S (L15 OR L20) NOT L21

FILE 'HCA' ENTERED AT 10:17:18 ON 17 AUG 2004

- L23 126935 S L19
- L24 7587 S L19/P
- L25 26196 S L22
- L26 47166 S L18

FILE 'LCA' ENTERED AT 10:18:34 ON 17 AUG 2004

FILE 'HCA' ENTERED AT 10:21:26 ON 17 AUG 2004

- L27 16554 S L4(2N)(L10 OR L11)
- L28 3181 S L27 AND (73/SX, SC OR 55/SX, SC OR 53/SX, SC)
- L29 13373 S L27 NOT L28
- L30 568 S L24 AND (L13 OR L7)
- L31 16503 S L29 OR L30 OR L7

FILE 'REGISTRY' ENTERED AT 10:25:48 ON 17 AUG 2004

FILE 'HCA' ENTERED AT 10:25:59 ON 17 AUG 2004

L32 TRA L31 1-4000 RN : 14182 TERMS

FILE 'REGISTRY' ENTERED AT 10:28:03 ON 17 AUG 2004 L33 14181 SEA L32

FILE 'HCA' ENTERED AT 10:29:28 ON 17 AUG 2004 L34 TRA L31 4001-10000 RN : 14825 TERMS

FILE 'REGISTRY' ENTERED AT 10:32:29 ON 17 AUG 2004 L35 14818 SEA L34

FILE 'HCA' ENTERED AT 10:33:30 ON 17 AUG 2004 L36 TRA L31 10001-16503 RN : 8822 TERMS

FILE 'REGISTRY' ENTERED AT 10:36:56 ON 17 AUG 2004 L37 8811 SEA L36

L38 32443 S L33 OR L35 OR L37

FILE 'LREGISTRY' ENTERED AT 10:44:06 ON 17 AUG 2004

- L39 STR
- L40 STR L39
- L41 STR

FILE 'REGISTRY' ENTERED AT 10:52:00 ON 17 AUG 2004 L42 50 S L40 OR L41 SSS SAM SUB=L38

```
L43
        7 S L41 SSS SAM SUB=L38
    FILE 'LREGISTRY' ENTERED AT 10:53:44 ON 17 AUG 2004
L44
               STR L40
               STR L44
L45
    FILE 'REGISTRY' ENTERED AT 10:58:14 ON 17 AUG 2004
L46
            50 S L45 OR L41 SSS SAM SUB=L38
    FILE 'LREGISTRY' ENTERED AT 10:59:10 ON 17 AUG 2004
L47
               STR L45
L48
               STR L45
    FILE 'REGISTRY' ENTERED AT 11:01:51 ON 17 AUG 2004
     11 S L47 OR L41 SSS SAM SUB=L38
L49
     FILE 'LREGISTRY' ENTERED AT 11:02:39 ON 17 AUG 2004
    FILE 'REGISTRY' ENTERED AT 11:09:08 ON 17 AUG 2004
L50
           50 S L45 OR L41 SSS SAM SUB=L38
L51
          4286 S L45 OR L41 SSS FULL SUB=L38
               SAVE L51 HRTZG243/A
L52
            11 S L47 OR L41 SSS SAM SUB=L51
L53
           257 S L47 OR L41 SSS FULL SUB=L51
               SAVE L51 HRTZG243A/A
               DELETE HRTZG243A/A
               SAVE L53 HRTZG243A/A
L54
          4029 S L51 NOT L53
   FILE 'HCA' ENTERED AT 11:13:18 ON 17 AUG 2004
L55
       207497 S L53
L56
       1996977 S L54
L57
            53 S L24 AND L55
L58
           674 S L24 AND L56
L59
        529810 S CARBONATE# OR ANHYDRIDE# OR EPOXIDE# OR POLYANHYDRIDE#
            29 S L57 AND L59
L60
             2 S L60 AND L13
L61
L62
           171 S L58 AND L59
L63
            15 S L62 AND L13
             2 S L62 AND L12
L64
L65
             1 S L57 AND L12
L66
        221878 S L4/TI
           93 S L62 AND L66
L67
        187433 S 49/SX,SC
L68
            64 S L62 AND L68
L69
L70
             6 S L69 AND L13
             1 S L69 AND L12
L71
            17 S L69 AND L10
L72
            17 S L72 AND L4
L73
L74
             QUE PREPARATION?
L75
            8 S L61 OR L64 OR L65 OR L70 OR L71
L76
            17 S L72 OR L73
L77
            27 S L76 OR L63
            29 S L77 OR L75
L78
L79
            26 S L78 AND 1907-2001/PY, PRY
            23 S L79 AND L74
L80
L81
            26 S L79 OR L80
L82
            26 S L81 AND L59
```

62 S L69 AND 1907-2000/PY, PRY L83 46 S L83 NOT L82 L84

FILE 'LCA' ENTERED AT 11:26:50 ON 17 AUG 2004

L85 6605 S RECOVER? OR WINN? OR PURIF? OR PURE# OR OXIDATION#

FILE 'HCA' ENTERED AT 11:34:29 ON 17 AUG 2004

L86 36536 S L6(3N)L85

29 S L84 AND L86 L87

L8825 S L78 AND 1907-2000/PY, PRY

L89 54 S L88 OR L88 OR L87 L90 54 S L89 AND (L55 OR L56)

FILE 'REGISTRY' ENTERED AT 11:38:40 ON 17 AUG 2004

=> d que stat L51

ARDITH.

I PERFORMED A TEXT SEARCH FOR MOLTEN SULFUR AND WHERE SULFUR WAS FORMED AS A PRODUCT AND TRANSFERED THE REGISTRY NUMBERS FROM THE CA RECORDS INTO THE REGISTRY FILE. I THEN DID A STRUCTURE SEARCH OF THE EXPOXIDE, ANHYDRIDE FROM THE REGISTRY NUMBERS THAT WERE TRANSFERRED FROM CHEMICAL ABSTRACTS INTO THE REGISTRY FILE.

10 SEA FILE=REGISTRY ABB=ON PLU=ON (108-32-7/BI OR 122-60-1/BI OR 123-54-6/BI OR 138-86-3/BI OR 2461-15-6/BI OR 25377-73-5/BI OR 26560-94-1/BI OR 26761-45-5/BI OR 7704-34-9/BI OR 7783-06-4/ 1899 SEA FILE=LCA ABB=ON PLU=ON SULFUR# OR SULPHUR# OR (SULFUR# OR SULPHUR#) (A) HYDRIDE## OR H2S OR MERCAPTO# OR SULFIDE# OR SULPHIDE# L5946220 SEA FILE=HCA ABB=ON PLU=ON (RETARDER? OR INHIBITOR? OR SUPPRESSOR? OR SCAVENG?) 1.6 666696 SEA FILE=HCA ABB=ON PLU=ON SULFUR# OR SULPHUR# OR (SULFUR# OR SULPHUR#) (A) HYDRIDE## OR HZS OR MERCAPTO# OR SULFIDE# OR SULPHIDE# T.7 3153 SEA FILE=HCA ABB=ON PLU=ON L5(3A)L6 L10

searched very broadly

5286 SEA FILE=LCA ABB=ON PLU=ON LIQUID# OR LIQ# OR MOLTEN# OR FLUID? OR MELTED? OR LIQUIF? OR COLLOIDAL#

3282 SEA FILE=LCA ABB=ON PLU=ON SUSPEN? OR DISPERS? OR COLLOID? L11OR EMULS? OR MICROEMULS? OR SLURR?

20487 SEA FILE=HCA ABB=ON PLU=ON L4(3A)(L10 OR L11) L13

6 SEA FILE=REGISTRY ABB=ON PLU=ON L2 AND 1-10/NR L15

PLU=ON L2 NOT L15 L16 4 SEA FILE=REGISTRY ABB=ON

L172 SEA FILE=REGISTRY ABB=ON PLU=ON L16 AND SULFUR

PLU=ON L17 AND 1-3/H L181 SEA FILE=REGISTRY ABB=ON

1 SEA FILE=REGISTRY ABB=ON PLU=ON L17 NOT L18 L19 L24

L27

7587 SEA FILE=HCA ABB=ON PLU=ON L19/P Preparation of Sulfur.
16554 SEA FILE=HCA ABB=ON PLU=ON L4(2A)(L10 OR L11)
3181 SEA FILE=HCA ABB=ON PLU=ON L27 AND (73/SX, SC OR 55/SX, SC OR 53/SY SC) L28 53/SX,SC)

I searched very broadly.

L29 13373 SEA FILE=HCA ABB=ON PLU=ON L27 NOT L28 L30 568 SEA FILE=HCA ABB=ON PLU=ON L24 AND (L13 OR L7) CHEMICAL 16503 SEA FILE=HCA ABB=ON PLU=ON L29 OR L30 OR L7 L31 ABSTRACT L32 TRANSFER PLU=ON L31 1-4000 RN: 14182 TERMS L33 14181 SEA FILE=REGISTRY ABB=ON PLU=ON L32 record L34 TRANSFER PLU=ON L31 4001-10000 RN: 14825 TERMS From WHICH 14818 SEA FILE=REGISTRY ABB=ON PLU=ON L34 L35 L36 TRANSEER PLU=ON L31 10001-16503 RN: 8822 TERMS REGISTRY # 3 8811 SEA FILE=REGISTRY ABB=ON L37 PLU=ON L36 L38 32443 SEA FILE=REGISTRY ABB=ON PLU=ON L33 OR L35 OR L37 WERETRANSFERE L41 INTO REC

seach was of these reg Is.

3 C C 2

Hertzog

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 3

STEREO ATTRIBUTES: NONE

VAR G1=8/13/18/26 NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 21

STEREO ATTRIBUTES: NONE

L51 4286 SEA FILE=REGISTRY SUB=L38 SSS FUL L45 OR L41

100.0% PROCESSED 17342 ITERATIONS

4286 ANSWERS

SEARCH TIME: 00.00.02

=> file hca

08/17/2004

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FILE COVERS 1907 - 13 Aug 2004 VOL 141 ISS 8 FILE LAST UPDATED: 13 Aug 2004 (20040813/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d L90 1-54 ti

- L90 ANSWER 1 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Difference detection methods using matched multiple dyes
- L90 ANSWER (2) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Hydrogen sulfide abatement in molten sulfur
- L90 ANSWER BOOF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Process for removing sulfur compounds from hydrocarbon streams
- L90 ANSWER 4/OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Method for oxidizing hydrogen sulfide to elemental sulfur
- L90 ANSWER (5) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Method and composition for removing CO2 and H2s from gas mixtures
- L90 ANSWER (6) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Study on removal of hydrogen sulfide from refinery acid gas by chemical absorption-catalytic oxidation in aqueous solution
- L90 ANSWER (7) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Recovery of sulfur from treated product gases from gasification of coal or heavy oils
- L90 ANSWER 8 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Processing of sulfur-containing materials from flue gas desulfurization or related sources for calcium compound recovery
- L90 ANSWER (9) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Process for preparing an agent based on colloidal sulfur

- obtained from sulfane for the protection of agricultural plants
- L90 ANSWER 10 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI A biological process for the reclamation of flue gas desulfurization gypsum using mixed sulfate-reducing bacteria with inexpensive carbon sources
- L90 ANSWER (11) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Manufacture of gypsum from sulfur derived from desulfurization of petroleum products
- L90 ANSWER 12 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Recycling of FGD gypsum to calcium carbonate and elemental sulfur using mixed sulfate-reducing bacteria with sewage digest as a carbon source
- L90 ANSWER (13) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Elemental sulfur recovery from flue gases.
- L90 ANSWER 14 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Combination of a sulfur compound and specific phosphorus compounds and their use in lubricating compositions, concentrates and greases
- L90 ANSWER (15) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Processing of residue from coal fluidized-bed gasification
- L90 ANSWER (16) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Method for recovering sulfur in a wet desulfurization of gases
- L90 ANSWER (17) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Recovery of technical sulfur from concentrates resulted from flotation of ores containing sulfur
- L90 ANSWER 18 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Flue Gas Desulfurization Scheme To Recover Elemental Sulfur
- L90 ANSWER (9) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Process for recovering sulfur from carbon dioxide- and hydrogen sulfide-containing acid gas mixtures
- L90 ANSWER 20 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Purification of sulfur
- L90 ANSWER 21 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Continuous biological process for the elimination of malodorous sulfur compounds present in gas mixtures
- L90 ANSWER (22) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Removal and recovery of sulfur dioxide from waste gases, especially from metallurgical plants, and from flue gases
- L90 ANSWER 23 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Petrology of Philippine geothermal systems and the application of alteration mineralogy to their assessment
- L90 ANSWER (24) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Manufacture of sublimed amorphous sulfur

- L90 ANSWER 25 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Soda recovery from metallurgical slags
- L90 ANSWER (26) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Production of water-suspendable liquid or solid sulfur concentrates
- L90 ANSWER (27) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI MHD seed recovery and regeneration
- L90 ANSWER(28)OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Process for removal of hydrogen sulfide from sour gas streams
- L90 ANSWER 29 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Method and apparatus for purifying raw gases from coal gasification while producing synthesis and fuel gas
- L90 ANSWER (30) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Effect of the method of copper oxide addition on chemisorption properties of zinc-containing sulfur-purification materials
- L90 ANSWER (31) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Process for removing hydrogen sulfide from gas streams
- L90 ANSWER 32 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Preparation of olefin adduct vulcanizing agents for rubbers
- L90 ANSWER (33) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Preparation of elementary sulfur from pyrites of any source and pyritic concentrates, in dry form at atmospheric pressure
- L90 ANSWER 34 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Method for purifying stratal and drainage water of sulfur pits
- L90 ANSWER 35 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI New gas scrubber removes H2S
- L90 ANSWER (36) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Method of winning of elemental sulfur from sulfur ore
- L90 ANSWER (37/OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Winning of pure sulfur from sulfur ore
- L90 ANSWER (38/OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Sulfur and calcium carbonate from gypsum
- L90 ANSWER (39/OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Desulfurization by three-stage combustion
- L90 ANSWER (40)OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Recovery of calcium carbonate and sulfur from FGD scrubber waste
- L90 ANSWER (41) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Removing hydrogen sulfide from gas streams
- L90 ANSWER (42) OF 54 HCA COPYRIGHT 2004 ACS on STN

- TI Conversion of alkali metal sulfate to the carbonate
- L90 ANSWER 43/OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Aqueous carbonate process design study
- L90 ANSWER 44 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Recovering calcium and sulfur compounds from a metallurgical waste material
- L90 ANSWER 45 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Rare anthropogenic and natural particles suspended in deep ocean waters
- L90 ANSWER 46 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Treatment of an alkali metal sulfide solution
- L90 ANSWER (47) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Removal of hydrogen sulfide from natural gas to obtain elemental sulfur
- L90 ANSWER (48) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Study of reagent conditions for autoclave melting of sulfur
- L90 ANSWER 49 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Recovery of sulfur dioxide
- L90 ANSWER 50 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Chemistry of the molten carbonate process for sulfur oxides removal stack gases
- L90 ANSWER 61/OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Treating liquid waste containing sulfur
- L90 ANSWER (52) OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Sulfur from Claus-process off-gases
- L90 ANSWER 53 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Converting soda slag from a blast furnace to useful products
- L90 ANSWER 54 OF 54 HCA COPYRIGHT 2004 ACS on STN
- TI Carbon oxide regenerant for **sulfur** recovery from **molten** salts
- => => d L90 2-7,9,11,13,15-20,22,24,26-28,30-31,33-34,36-54 bib abs hitind hitstr
- L90 ANSWER 2 OF 54 HCA COPYRIGHT 2004 ACS on STN
- 136:312044 Hydrogen sulfide abatement in molten sulfur. Schield, John A.; Weers, Jerry J.; Cappel, Weldon J.; Roof, Glenn L. (Baker Hughes Incorporated, USA). Eur. Pat. Appl. EP 1197473 A1 20020417, 15 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO. (English). CODEN: EPXXDW. APPLICATION: EP 2001-308773 20011015. PRIORITY: US 2000-PV240140 20001013; US 2001-975438 20011009.
- The present invention provides a method for inhibiting the evolution of H2S from sulfhydryl compds. in molten sulfur by using scavenging agents such as anhydrides and polymers thereof, conjugated ketones, carbonates, epoxides, monoesters and diesters of unsatd. dicarboxylic acids and polymers of these esters, and the like and mixts. thereof. In one

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embodiment, it is preferred that the scavenging agent is in liquid
form at contact temperature with the molten sulfur. In
another embodiment, the scavenging agent may be atomized into the vapor
space over the molten sulfur to contact the
sulfur with the agent.
ICM C01B017-02
49-1 (Industrial Inorganic Chemicals)
hydrogen sulfide abatement molten sulfur
Naphthenic acids, uses
RL: MOA (Modifier or additive use); TEM (Technical or engineered material
use); USES (Uses)
   (copper salts, scavenging agent; hydrogen sulfide abatement
   in molten sulfur)
Carboxylic acids, uses
RL: TEM (Technical or engineered material use); USES (Uses)
   (dicarboxylic, diesters, scavenging agent; hydrogen sulfide
   abatement in molten sulfur)
Carboxylic acids, uses
RL: TEM (Technical or engineered material use); USES (Uses)
   (esters, scavenging agent; hydrogen sulfide abatement in
   molten sulfur)
Mercapto compounds (inorganic)
RL: MSC (Miscellaneous)
   (hydrogen sulfide abatement in molten
   sulfur)
Naphthenic acids, uses
RL: MOA (Modifier or additive use); TEM (Technical or engineered material
use); USES (Uses)
   (iron salts, scavenging agent; hydrogen sulfide abatement in
   molten sulfur)
Fatty acids, uses
RL: MOA (Modifier or additive use); TEM (Technical or engineered material
use); USES (Uses)
   (linseed-oil, epoxidized, Me esters, scavenging agent; hydrogen
   sulfide abatement in molten sulfur)
Alcohols, uses
RL: TEM (Technical or engineered material use); USES (Uses)
   (polyhydric, scavenging agent; hydrogen sulfide abatement in
   molten sulfur)
Anhydrides
  Carbonates, uses
  Epoxides
  Polyanhydrides
RL: TEM (Technical or engineered material use); USES (Uses)
   (scavenging agent; hydrogen sulfide abatement in
   molten sulfur)
Fatty acids, uses
RL: MOA (Modifier or additive use); TEM (Technical or engineered material
use); USES (Uses)
   (soya, epoxidized, Me esters, VikoFlex 7010, scavenging agent; hydrogen
   sulfide abatement in molten sulfur)
Ketones, uses
RL: TEM (Technical or engineered material use); USES (Uses)
   (unsatd., conjugated, scavenging agent; hydrogen sulfide
   abatement in molten sulfur)
Naphthenic acids, uses
RL: MOA (Modifier or additive use); TEM (Technical or engineered material
use); USES (Uses)
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(zinc salts, scavenging agent; hydrogen sulfide abatement in
        molten sulfur)
IT
     26761-45-5
     RL: MOA (Modifier or additive use); TEM (Technical or engineered material
     use); USES (Uses)
        (Epoxide 248, scavenging agent; hydrogen sulfide
        abatement in molten sulfur)
IT
     122-60-1, Heloxy 63
     RL: MOA (Modifier or additive use); TEM (Technical or engineered material
     use); USES (Uses)
        (Heloxy 63, scavenging agent; hydrogen sulfide abatement in
        molten sulfur)
     7704-34-9P, Sulfur, preparation
IT
     RL: PUR (Purification or recovery); PREP (Preparation)
        (hydrogen sulfide abatement in molten
        sulfur)
     7783-06-4, Hydrogen sulfide, processes
IT
     RL: REM (Removal or disposal); PROC (Process)
        (hydrogen sulfide abatement in molten
IT
     108-32-7, Propylene carbonate 123-54-6,
     2,4-Pentanedione, uses 138-86-3, Dipentene 2461-15-6,
     2-Ethylhexylglycidyl ether 25377-73-5, Dodecenylsuccinic
     anhydride 26560-94-1
     RL: MOA (Modifier or additive use); TEM (Technical or engineered material
     use); USES (Uses)
        (scavenging agent; hydrogen sulfide abatement in
        molten sulfur)
     26761-45-5
IT
     RL: MOA (Modifier or additive use); TEM (Technical or engineered material
     use); USES (Uses)
        (Epoxide 248, scavenging agent; hydrogen sulfide
        abatement in molten sulfur)
RN
     26761-45-5 HCA
     Neodecanoic acid, oxiranylmethyl ester (9CI) (CA INDEX NAME)
CN
     CH2-O-C-(C9H19-neo)
     122-60-1, Heloxy 63
IT
     RL: MOA (Modifier or additive use); TEM (Technical or engineered material
     use); USES (Uses)
        (Heloxy 63, scavenging agent; hydrogen sulfide abatement in
        molten sulfur)
RN
     122-60-1 HCA
CN
     Oxirane, (phenoxymethyl) - (9CI) (CA INDEX NAME)
     CH2-OPh
ΙT
     7704-34-9P, Sulfur, preparation
     RL: PUR (Purification or recovery); PREP (Preparation)
        (hydrogen sulfide abatement in molten
        sulfur)
```

RN 7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

1T 108-32-7, Propylene carbonate 123-54-6,
2,4-Pentanedione, uses 2461-15-6, 2-Ethylhexylglycidyl ether
25377-73-5, Dodecenylsuccinic anhydride
26560-94-1

RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)

(scavenging agent; hydrogen sulfide abatement in

molten sulfur)

RN 108-32-7 HCA

CN 1,3-Dioxolan-2-one, 4-methyl- (9CI) (CA INDEX NAME)

RN 123-54-6 HCA

CN 2,4-Pentanedione (8CI, 9CI) (CA INDEX NAME)

RN 2461-15-6 HCA

CN Oxirane, [[(2-ethylhexyl)oxy]methyl]- (9CI) (CA INDEX NAME)

$$\overset{\text{O}}{\underset{\text{CH}_2\text{- O- CH}_2\text{- CH- Bu-n}}{\text{Et}}}$$

RN 25377-73-5 HCA

CN 2,5-Furandione, 3-(dodecenyl)dihydro- (9CI) (CA INDEX NAME)

CM 1

CRN 2561-85-5 CMF C16 H28 O3

RN 26560-94-1 HCA

CN 2-Butenedioic acid (2Z)-, mono(2-hydroxyethyl) ester (9CI) (CA INDEX NAME)

Double bond geometry as shown.

L90 ANSWER 3 OF 54 HCA COPYRIGHT 2004 ACS on STN

135:305979 Process for removing sulfur compounds from hydrocarbon streams.

Pittman, Rusty; Arena, Blaise J.; Janssen, Albert J. (UOP LLC, USA). U.S.

US 6306288 B1 20011023, 6 pp., Cont.-in-part of U.S. Ser. No. 61,661,
abandoned. (English). CODEN: USXXAM. APPLICATION: US 1999-426818
19991022. PRIORITY: US 1998-61661 19980417.

AB A process for removing H2S and mercaptans from a hydrocarbon stream is disclosed. A hydrocarbon stream such as a LPG stream is contacted with a weakly basic stream, e.g., a Na bicarbonate stream to extract the H2S and mercaptans from the hydrocarbon stream into the basic stream. The basic stream is now treated in a reactor containing a sulfide-oxidizing microorganism to convert the H2S to S and the mercaptans to disulfides. Finally, the S and disulfides are separated from the basic aqueous stream which can be recycled and used to treat a fresh hydrocarbon stream. The treated hydrocarbon stream is purified to the point that it passes the Cu strip test, while the purified basic stream contains <0.08 g S/L.

IC ICM C10G019-08

ICS C10G019-00; C10G032-00

NCL 208235000

CC 51-4 (Fossil Fuels, Derivatives, and Related Products)
 Section cross-reference(s): 49

IT Disulfides

RL: PUR (Purification or recovery); PREP (Preparation) (recovery in removing of sulfur compds. from hydrocarbon streams)

IT 144-55-8, Sodium bicarbonate, uses 497-19-8, Sodium carbonate, uses 1336-21-6, Ammonium hydroxide RL: TEM (Technical or engineered material use); USES (Uses) (in removing of sulfur compds. from hydrocarbon streams)

7704-34-9P, Sulfur, preparation

RL: PUR (Purification or recovery); PREP (Preparation)
 (recovery in removing of sulfur compds. from
 hydrocarbon streams)

IT 144-55-8, Sodium bicarbonate, uses 497-19-8, Sodium carbonate, uses

RL: TEM (Technical or engineered material use); USES (Uses) (in removing of sulfur compds. from hydrocarbon streams)

RN 144-55-8 HCA

CN Carbonic acid monosodium salt (8CI, 9CI) (CA INDEX NAME)

О || но- с- он

Na

RN 497-19-8 HCA

CN Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)

НО— С— ОН

●2 Na

RN 7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

L90 ANSWER 4 OF 54 HCA COPYRIGHT 2004 ACS on STN

135:290881 Method for oxidizing hydrogen sulfide to elemental sulfur. Li,
Kuo-tseng; Chi, Ren-hai (National Science Council, Taiwan). U.S. US
6299851 B1 20011009, 7 pp., Cont.-in-part of U.S. Ser. No. 277,301.
(English). CODEN: USXXAM. APPLICATION: US 1999-334851 19990617.
PRIORITY: US 1999-277301 19990326.

AB A method for selectively oxidizing hydrogen sulfide to elemental sulfur is disclosed. The method is performed at 50°to 400° and at a pressure ranged from 0.1 to 50 atmospheric. The elemental sulfur can be effectively recovered from a gas mixture containing hydrogen sulfide in the presence of a catalyst. The catalyst includes a vanadium-containing material and a catalytic substance selected from the group consisting of scandium (Sc), yttrium (Y), lanthanum (La), samarium (Sm) and compds. thereof. In another embodiment, this catalyst further includes an antimony-containing promoter (antimony compds.) which further exhibit a more effective catalytic performance.

IC ICM C01B017-04

NCL 423573100

CC 49-10 (Industrial Inorganic Chemicals)

ST sulfur manuf hydrogen sulfide oxidn catalyst

69-72-7D, Salicylic acid, vanadium complexes 1314-23-4, IT Zirconia, uses 1327-33-9, Antimony oxide 1344-28-1, Alumina, uses 6218-52-6, Vanadium formate 7439-91-0, Lanthanum, uses 7440-19-9, Samarium, uses 7440-20-2, Scandium, uses 7440-36-0, Antimony, uses 7440-45-1, Cerium, uses 7440-62-2, Vanadium, uses 7440-65-5, Yttrium, uses 7631-86-9, Silica, uses 11099-11-9, Vanadium oxide 11115-67-6, Ammonium vanadate 11130-24-8, Vanadium sulfide 12070-10-9, Vanadium 12627-52-0, Antimony sulfide 12674-04-3, Vanadium nitride 12713-06-3, Vanadium hydride 13463-67-7, Titania, uses 14 Vanadium phosphate **14974-48-2**, Vanadium oxalate 16785-81-2, 14542-94-0, Vanadium sulfate 20644-87-5, Vanadium carbonyl 24492-29-3, Vanadium 39318-26-8, Chromium vanadium oxide oleate 37353-31-4, Vanadate 39349-74-1, Antimonate 51891-70-4, Vanadium carbonate 57348-87-5, Vanadium phosphide 63465-09-8, Vanadium acetate 67422-42-8, Antimony carbide 77414-04-1 102857-58-9, Vanadium 123782-01-4 132036-01-2 152761-81-4, Antimony hydroxide hydroxide

259744-19-9 364427-27-0

RL: CAT (Catalyst use); USES (Uses)

(method for oxidizing hydrogen sulfide to elemental sulfur)

IT 7704-34-9P, Sulfur, preparation

RL: IMF (Industrial manufacture); PEP (Physical, engineering or chemical process); PREP (Preparation); PROC (Process)

(method for oxidizing hydrogen sulfide to elemental sulfur)

IT 69-72-7D, Salicylic acid, vanadium complexes 14974-48-2,

Vanadium oxalate

RL: CAT (Catalyst use); USES (Uses)

(method for oxidizing hydrogen sulfide to elemental sulfur)

RN 69-72-7 HCA

CN Benzoic acid, 2-hydroxy- (9CI) (CA INDEX NAME)

RN 14974-48-2 HCA

CN Ethanedioic acid, vanadium salt (9CI) (CA INDEX NAME)

\bullet x V(x)

IT 7704-34-9P, Sulfur, preparation

RL: IMF (Industrial manufacture); PEP (Physical, engineering or chemical process); PREP (Preparation); PROC (Process)

(method for oxidizing hydrogen sulfide to elemental sulfur)

RN 7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

L90 ANSWER 5 OF 54 HCA COPYRIGHT 2004 ACS on STN

134:254262 Method and composition for removing CO2 and H2S from gas mixtures. Chao, I-Meen (Eickmeyer & Associates, USA). PCT Int. Appl. WO 2001019496 A1 20010322, 24 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO,

LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA,

ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH,

CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE,

NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO

2000-US25397 20000915. PRIORITY: US 1999-PV154009 19990915.

AB Process is disclosed for removal of CO2 and H2S from a sour gas mixture (e.g., natural gas, Fe ore reduction gas). The process uses a continuous absorption process with an alkaline aqueous liquid mixture containing

Hertzog 10/688,243 08/17/2004

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15-40 weight% of equivalent K2CO3 and is conducted at 150-300°F. The
     H2S is removed from the sour gas as elementary S. CO2 is stripped
     from the aqueous alkaline solution and vented directly into the atmospheric
since the S that
     usually contaminates the CO2 was removed earlier in the process as
     elementary S.
TC.
     ICM B01D053-14
     ICS B01D053-52
     48-1 (Unit Operations and Processes)
     Section cross-reference(s): 49, 51
     carbon dioxide removal gas mixt absorption; hydrogen sulfide
     removal gas mixt absorption; natural gas sweetening absorption
     Absorption
        (for removal of carbon dioxide and hydrogen sulfide from gas
        mixts.)
IT
     Natural gas, processes
     RL: PEP (Physical, engineering or chemical process); PUR (Purification or
     recovery); PREP (Preparation); PROC (Process)
        (removal of carbon dioxide and hydrogen sulfide from gas
        mixts. by absorption)
ΙT
     584-08-7, Potassium carbonate 1314-62-1, Vanadium
     oxide (V2O5), uses 10043-35-3, Boric acid (H3BO3), uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (in absorption solution for removal of carbon dioxide and hydrogen
        sulfide from gas mixts.)
IT
     7704-34-9P, Sulfur, preparation
     RL: IMF (Industrial manufacture); PREP (Preparation)
        (recovery in removal of carbon dioxide and hydrogen sulfide
        from gas mixts. by absorption)
     124-38-9, Carbon dioxide, processes 7783-06-4, Hydrogen
     sulfide, processes
     RL: REM (Removal or disposal); PROC (Process)
        (removal from gas mixture by absorption)
     584-08-7, Potassium carbonate
     RL: TEM (Technical or engineered material use); USES (Uses)
        (in absorption solution for removal of carbon dioxide and hydrogen
        sulfide from gas mixts.)
RN
     584-08-7 HCA
CN
     Carbonic acid, dipotassium salt (8CI, 9CI) (CA INDEX NAME)
   \circ
HO- C- OH
 ●2 K
IT
     7704-34-9P, Sulfur, preparation
     RL: IMF (Industrial manufacture); PREP (Preparation)
        (recovery in removal of carbon dioxide and hydrogen sulfide
        from gas mixts. by absorption)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
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L90 ANSWER 6 OF 54 HCA COPYRIGHT 2004 ACS on STN

133:270762 Study on removal of hydrogen sulfide from refinery acid
gas by chemical absorption-catalytic oxidation in aqueous solution. Dong,
Qun; Wu, Xianchun; Shan, Xilin; Wu, Guanjing; Zhang, Jianguo (Daqing
Petroleum Institute, Anda, 151400, Peop. Rep. China). Shiyou Lianzhi Yu
Huagong, 31(9), 17-19 (Chinese) 2000. CODEN: SLYHEE. ISSN:
1005-2399. Publisher: Shiyou Lianzhi Yu Huagong Zazhishe.

AB A process for removal of hydrogen sulfide from refinery acid gas by wet chemical absorption/catalytic oxidation in a basic solution containing catalyst

was conducted in the laboratory The effects of flow rate ratio of gas/liquid in the absorber, the flow rate of air and the residence time of the basic solution in the regeneration tower on the performance of hydrogen sulfide removal from refinery acid gas and elemental S recovery were studied. The test result showed that the said process is expected to be used in refineries for acid gas treatment.

CC 59-4 (Air Pollution and Industrial Hygiene) Section cross-reference(s): **49**, 51, 60

ST hydrogen **sulfide** removal refinery acid gas absorption catalytic oxidn; petroleum refinery acid gas wet absorption oxidn **sulfur** recovery

IT Petroleum products

(refinery gases, acid; hydrogen **sulfide** removal from refinery acid gas by wet chemical absorption-catalytic oxidation and **sulfur** recovery)

IT 497-19-8, Sodium carbonate, processes

RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(absorption by; hydrogen **sulfide** removal from refinery acid gas by wet chemical absorption-catalytic oxidation and **sulfur** recovery)

IT 7772-98-7P, Sodium thiosulfate

RL: BYP (Byproduct); PREP (Preparation)

(hydrogen **sulfide** removal from refinery acid gas by wet chemical absorption-catalytic oxidation and **sulfur** recovery)

IT 7704-34-9P, Sulfur, preparation

RL: PUR (Purification or recovery); PREP (Preparation)
(hydrogen sulfide removal from refinery acid gas by wet chemical absorption-catalytic oxidation and sulfur recovery)

IT 7783-06-4, Hydrogen sulfide, processes

RL: REM (Removal or disposal); PROC (Process)

(hydrogen sulfide removal from refinery acid gas by wet chemical absorption-catalytic oxidation and sulfur recovery)

IT 497-19-8, Sodium carbonate, processes

RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(absorption by; hydrogen sulfide removal from refinery acid

gas by wet chemical absorption-catalytic oxidation and sulfur recovery)

RN 497-19-8 HCA

CN Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)

О || НО— С— ОН

●2 Na

IT 7704-34-9P, Sulfur, preparation

RL: PUR (Purification or recovery); PREP (Preparation) (hydrogen sulfide removal from refinery acid gas by wet chemical absorption-catalytic oxidation and sulfur recovery)

RN 7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

L90 ANSWER 7 OF 54 HCA COPYRIGHT 2004 ACS on STN

132:37943 Recovery of sulfur from treated product gases from gasification of coal or heavy oils. Yoshida, Kunikatsu; Yamada, Michio; Ueda, Akio; Ueda, Toshiyuki (Babcock-Hitachi K. K., Japan). Jpn. Kokai Tokkyo Koho JP 11347348 A2 19991221 Heisei, 4 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1998-154006 19980603.

AB Raw product gases containing H2S from gasification of coal or heavy oils are treated by wet scrubbing with water to sep. dust, oxidation of the treated gases to convert H2S into SO2, contacting the gases with an alkali slurry containing CaCO3, and further oxidation with divalent metal ions (especially,

Fe2+ or

Co2+) to convert sulfites into gypsum, and then recovering elemental S (as gypsum) from the oxidized slurry. In one embodiment, the waste waters containing divalent metal ions (especially, Fe2+ or Co2+) from dust during wet scrubbing can be utilized for increasing the oxidation of CaSO3 into gypsum.

IC ICM B01D053-50 ICS B01D053-77

CC 51-20 (Fossil Fuels, Derivatives, and Related Products)
Section cross-reference(s): 49

sulfur recovery raw product coal gasification; gypsum coal gasification raw product desulfurization

IT Limestone, uses

RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(alkali scrubbing solution containing; recovery of sulfur from treated product gases from gasification of coal or heavy oils)

IT Fuel oil

(heavy; recovery of sulfur from treated product gases from gasification of coal or heavy oils)

IT Coal gasification

(recovery of sulfur from treated product gases from gasification of coal or heavy oils)

IT 471-34-1, Calcium carbonate, uses

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RL: PEP (Physical, engineering or chemical process); TEM (Technical or
     engineered material use); PROC (Process); USES (Uses)
        (alkali scrubbing solution containing; recovery of sulfur
        from treated product gases from gasification of coal or heavy oils)
ΙT
     15438-31-0, Iron ion (Fe2+), uses 22541-53-3, Cobalt ion (Co2+), uses
     RL: PEP (Physical, engineering or chemical process); TEM (Technical or
     engineered material use); PROC (Process); USES (Uses)
        (oxidation by; recovery of sulfur from
        treated product gases from gasification of coal or heavy oils)
                                      13397-24-5P, Gypsum,
IT
     7704-34-9P, Sulfur, preparation
     preparation
     RL: BYP (Byproduct); PREP (Preparation)
        (recovery of sulfur from treated product gases from
        gasification of coal or heavy oils)
ΙT
     471-34-1, Calcium carbonate, uses
     RL: PEP (Physical, engineering or chemical process); TEM (Technical or
     engineered material use); PROC (Process); USES (Uses)
        (alkali scrubbing solution containing; recovery of sulfur
        from treated product gases from gasification of coal or heavy oils)
RN
     471-34-1 HCA
CN
     Carbonic acid calcium salt (1:1) (8CI, 9CI) (CA INDEX NAME)
HO-C-OH
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Ca

S

L90 ANSWER 9 OF 54 HCA COPYRIGHT 2004 ACS on STN

128:85437 Process for preparing an agent based on colloidal

sulfur obtained from sulfane for the protection of agricultural
plants. Koran, Jiri; Bouska, Petr; Buryan, Petr; Konas, Jiri; Kratochvil,
Jiri; Pospisil, Jaroslav; Zacher, Jan (Spolana, Czech Rep.). Czech Rep.

CZ 282157 B6 19970514, 4 pp. (Czech). CODEN: CZXXED.

APPLICATION: CZ 1990-6423 19901220.

A new process for the preparation of colloid sulfur.

As a new process for the preparation of colloid sulfur fungicides is described. The process eliminates costly energy—and machinery—demanding milling, mixing, and drying associated with uneven sulfur particle size distribution. The process involves formation of a suspension from gaseous sulfan and aqueous solns. of iron complexes. Sulfan (4-6 g/m3) is oxidized to sulfur by the complex formed from FeSO4.7H2O, Na EDTA, H2SO4, and Na2CO3. The sulfur particle size distribution is usually sufficient without further operations. If needed, the resulting mix is milled, sieved (sieve pore size 0.18 x 0.1 mm), and dried in a spray dryer

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at 110-140°C. The residues of iron complex solution remaining in the colloid sulfur preparation do not need to be removed as they are harmless in the soil/plant environment and may even contribute some nutrients. The suspension containing 40-99.5~mass% sulfur is mixed with 1-55~mass%mass% of a surfactant, 0.1-8 mass% NH4HCO3, and 1-20 mass% of aqueous ammonia (concentration 1-25%). The colloid sulfur can be mixed with sulfite solution (surfactant) and ground sulfur (with 3.9% mineral oil) in the preparation of final product. The process can efficiently utilize sulfur byproducts from gas desulfurization. A01N059-02; A01N025-04; C01B017-10 5-1 (Agrochemical Bioregulators) colloid sulfur prepn sulfan iron complex; fungicide

colloid sulfur prepn sulfan iron

ΙT Fungicides

> (colloidal sulfur preparation from sulfane for the protection of agricultural plants)

ΙT 7704-34-9P, Sulfur, preparation

RL: IMF (Industrial manufacture); PREP (Preparation) (colloidal sulfur preparation from sulfane and iron complex for fungicide production)

ΙT 139-33-3 497-19-8, Sodium carbonate, reactions 1066-33-7, Ammonium bicarbonate 7439-89-6D, Iron, complexes, 7664-41-7, Ammonia, reactions 7664-93-9, Sulfuric acid, reactions 7782-63-0, Ferrous sulfate heptahydrate 14265-45-3, Sulfite reactions

RL: RCT (Reactant); RACT (Reactant or reagent) (colloidal sulfur preparation from sulfane for the protection of agricultural plants)

IT7704-34-9P, Sulfur, preparation

37331-50-3, Sulfane

RL: IMF (Industrial manufacture); PREP (Preparation) (colloidal sulfur preparation from sulfane and iron complex for fungicide production)

7704-34-9 HCA RN

Sulfur (8CI, 9CI) (CA INDEX NAME)

S

TΤ 139-33-3 497-19-8, Sodium carbonate, reactions 1066-33-7, Ammonium bicarbonate RL: RCT (Reactant); RACT (Reactant or reagent) (colloidal sulfur preparation from sulfane for the protection of agricultural plants)

RN 139-33-3 HCA

Glycine, N, N'-1, 2-ethanediylbis[N-(carboxymethyl)-, disodium salt (9CI) CN (CA INDEX NAME)

●2 Na

497-19-8 HCA

Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)

О || но— с— он

●2 Na

RN 1066-33-7 HCA

CN Carbonic acid, monoammonium salt (8CI, 9CI) (CA INDEX NAME)

О || НО— С— ОН

NH3

L90 ANSWER 11 OF 54 HCA COPYRIGHT 2004 ACS on STN

126:33989 Manufacture of gypsum from sulfur derived from desulfurization of petroleum products. Shiozawa, Suguru (Kowa Co, Japan).

Jpn. Kokai Tokkyo Koho JP 08283018 A2 19961029 Heisei, 3 pp.
(Japanese). CODEN: JKXXAF. APPLICATION: JP 1995-110234 19950412.

- AB The process comprises burning molten S formed in the desulfurization to give SO2, burning the SO2 with catalysts to give SO3, converting the SO3 to H2SO4 by absorption with H2SO4, treating the H2SO4 with a CaCO3 slurry to give a CaSO4 slurry, and dewatering the slurry. The byproduct S is utilized, and the gypsum obtained is suitable for cement or gypsum boards.
- IC ICM C01F011-46
- CC 49-5 (Industrial Inorganic Chemicals)
 Section cross-reference(s): 51, 58
- ST gypsum manuf petroleum desulfurization sulfur recycling; calcium sulfate manuf petroleum desulfurization sulfur
- IT Petroleum refining

(desulfurization; gypsum manufacture from sulfur derived from desulfurization of petroleum products)

IT Recycling

(gypsum manufacture from **sulfur** derived from desulfurization of petroleum products)

IT 7704-34-9P, Sulfur, preparation

RL: BYP (Byproduct); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)

(gypsum manufacture from **sulfur** derived from desulfurization of petroleum products)

IT 7778-18-9P, Calcium sulfate

RL: IMF (Industrial manufacture); PEP (Physical, engineering or chemical process); PREP (Preparation); PROC (Process)

(gypsum manufacture from **sulfur** derived from desulfurization of petroleum products)

TT 7446-09-5P, Sulfur dioxide, preparation 7446-11-9P,
Sulfur trioxide, preparation 7664-93-9P, Sulfuric acid,
preparation

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RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT
     (Reactant or reagent)
        (gypsum manufacture from sulfur derived from desulfurization of
        petroleum products)
IT
     13397-24-5P, Gypsum, preparation
     RL: IMF (Industrial manufacture); TEM (Technical or engineered material
     use); PREP (Preparation); USES (Uses)
        (gypsum manufacture from sulfur derived from desulfurization of
        petroleum products)
IT
     471-34-1, Calcium carbonate, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (gypsum manufacture from sulfur derived from desulfurization of
        petroleum products)
IT
     7704-34-9P, Sulfur, preparation
     RL: BYP (Byproduct); RCT (Reactant); PREP (Preparation); RACT (Reactant or
        (gypsum manufacture from sulfur derived from desulfurization of
        petroleum products)
     7704-34-9 HCA
RN
CN
     Sulfur (8CI, 9CI)
                       (CA INDEX NAME)
S
IT
     471-34-1, Calcium carbonate, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (gypsum manufacture from sulfur derived from desulfurization of
        petroleum products)
RN
     471-34-1 HCA
CN
     Carbonic acid calcium salt (1:1) (8CI, 9CI) (CA INDEX NAME)
   0
HO-C-OH
```

● Ca

L90 ANSWER 13 OF 54 HCA COPYRIGHT 2004 ACS on STN 124:236405 Elemental sulfur recovery from flue gases.. Kerti, Jozsef; Kerti, Zsolt (Hung.). Hung. Teljes HU 70675 A2 19951030, 9 pp. (Hungarian). CODEN: HUXXBU. APPLICATION: HU 1992-9203160 19921006. Flue gases from thermal power stations, containing SO2, are contacted with CaS, CaCO3, Na2S, or/and Na2CO3 deposited on a porous support at 300-1000° in a reactor. The resulting sulfites are reduced with C, CO, H, and/or natural gas and the elemental S reaction product is condensed in a in a sept. chamber. IC ICM B01D053-34 49-2 (Industrial Inorganic Chemicals) CC Section cross-reference(s): 59 STsulfur recovery flue gas IT Flue gases Recycling (recovery from thermal power station flue gases, containing SO2, by

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treating with sulfides and/or carbonates and reducing)
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (recovery from thermal power station flue gases, containing SO2, by
        treating with sulfides and/or carbonates and reducing)
IT
     471-34-1, Calcium carbonate, uses 497-19-8,
     Disodium carbonate, uses 630-08-0, Carbon monoxide, uses
     1313-82-2, Sodium sulfide, uses 1333-74-0, Hydrogen, uses
     7440-44-0, Carbon, uses 20548-54-3, Calcium sulfide
     RL: NUU (Other use, unclassified); USES (Uses)
        (recovery from thermal power station flue gases, containing SO2,
        by treating with sulfides and/or carbonates and reducing)
IT
     7446-09-5, Sulfur dioxide, processes
     RL: PEP (Physical, engineering or chemical process); PROC (Process)
        (recovery from thermal power station flue gases, containing SO2,
        by treating with sulfides and/or carbonates and reducing)
IT
     7704-34-9P, Sulfur, preparation
     RL: PUR (Purification or recovery); PREP (Preparation)
        (recovery from thermal power station flue gases, containing SO2,
        by treating with sulfides and/or carbonates and reducing)
ΙT
     471-34-1, Calcium carbonate, uses 497-19-8,
     Disodium carbonate, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (recovery from thermal power station flue gases, containing SO2,
        by treating with sulfides and/or carbonates and reducing)
RN
     Carbonic acid calcium salt (1:1) (8CI, 9CI) (CA INDEX NAME)
CN
   O
HO-C-OH
 Ca
RN
     497-19-8 HCA
CN
     Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)
HO-C-OH
•2 Na
     7704-34-9P, Sulfur, preparation
     RL: PUR (Purification or recovery); PREP (Preparation)
        (recovery from thermal power station flue gases, containing SO2,
        by treating with sulfides and/or carbonates and reducing)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
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L90 ANSWER 15 OF 54 HCA COPYRIGHT 2004 ACS on STN
123:321351 Processing of residue from coal fluidized-bed
     gasification. Herbert, Peter; Reimert, Rainer; Stroeder, Michael
     (Metallgesellschaft AG, Germany). Ger. Offen. DE 4410598 A1
     19950928, 6 pp. (German). CODEN: GWXXBX. APPLICATION: DE
     1994-4410598 19940326.
     A residue (containing coke 8-80, alkaline earth metal sulfide (e.g.,
     CaS) 2-45, unreacted alkaline earth metal oxide (e.g., CaO) 1-25 weight%, and
ash
     balance) from fluidized-bed gasification of S-rich coal at
     700-1100° in the presence of an alkaline earth metal carbonate
     or alkaline earth metal oxide for (partial) desulfurization is withdrawn from
     the reactor, optionally milled, and mixed with an acid-containing aqueous
solution
     (e.g., CO2, H2SO4). After removal of liberated H2S (which is
     used for S manufacture), the residue containing ash, coke, and alkaline earth
     salt (e.g., CaCO3, CaSO4) is optionally milled and charged into a
     flotation zone to sep. coke (which is recycled to gasification) and the
     remaining residue containing ash and alkaline earth metal salt. The residue is
     dewatered in filter presses, rinsed, filtered again, and disposed.
     ICM B01D053-34
     60-5 (Waste Treatment and Disposal)
     Section cross-reference(s): 49, 51
     coal gasification residue processing; coke recycling gasification residue;
     sulfur manuf gasification residue
IT
     Coal gasification
        (fluidized-bed, processing of residue from)
TТ
     124-38-9, Carbon dioxide, reactions 7664-93-9, Sulfuric acid,
     reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (in removal of alkaline earth metal sulfides from coal
        gasification residue)
IT
     7704-34-9P, Sulfur, preparation
     RL: IMF (Industrial manufacture); PREP (Preparation)
        (manufacture of sulfur from hydrogen sulfide liberated
        during coal gasification residue processing)
IT
     471-34-1, Calcium carbonate (CaCO3), uses 7778-18-9,
     Calcium sulfate (CaSO4)
     RL: NUU (Other use, unclassified); USES (Uses)
        (processing and disposal of coal gasification residue containing)
IT
     124-38-9, Carbon dioxide, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
```

124-38-9 HCA

Carbon dioxide (8CI, 9CI) (CA INDEX NAME) CN

gasification residue)

0 = c = 0

RN

IT 7704-34-9P, Sulfur, preparation RL: IMF (Industrial manufacture); PREP (Preparation) (manufacture of sulfur from hydrogen sulfide liberated during coal gasification residue processing) RN 7704-34-9 HCA

(in removal of alkaline earth metal sulfides from coal

```
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
     471-34-1, Calcium carbonate (CaCO3), uses
IT
     RL: NUU (Other use, unclassified); USES (Uses)
         (processing and disposal of coal gasification residue containing)
RN
     471-34-1 HCA
CN
     Carbonic acid calcium salt (1:1) (8CI, 9CI) (CA INDEX NAME)
    \circ
HO-C-OH
  ● Ca
L90 ANSWER 16 OF 54 HCA COPYRIGHT 2004 ACS on STN
123:117472 Method for recovering sulfur in a wet desulfurization of gases.
     Kowalik, Waldemar; Demusiak, Grzegorz (Instytut Gornictwa Naftowego i
     Gazownictwa, Pol.). Pol. PL 163962 Bl 19940531, 4 pp.
     (Polish). CODEN: POXXA7. APPLICATION: PL 1990-287799 19901115.
     The process includes absorbing sulfur contaminants in a solution containing
AΒ
     hydroquinone and sodium carbonate, regenerating the absorption
     solution by aeration, sedimentation of a sepd.sulfur foam, and filtering
     unpurified sulfur suspension, followed by repulping
     sulfur paste, washing, and filtering the resulting purified
     sulfur suspension, from which flotation sulfur
     is obtained. The absorption is performed at a hydroquinone concentration of
     0.4-2.0 g/L, preferably 0.6-1.2 g/L, the regeneration is done at an air
     volume flow rate/absorption solution flow rate of (2-10):1, preferably (3-7):1,
     and the filtration of unpurified and purified sulfur
     suspensions is performed at 288-340 K.
IC
     ICM C01B017-05
     ICS B01D053-14
CC
     49-1 (Industrial Inorganic Chemicals)
     Section cross-reference(s): 51
TΤ
     123-31-9, Hydroquinone, uses 497-19-8, Sodium carbonate
     RL: NUU (Other use, unclassified); USES (Uses)
        (absorption solution component; in recovery of sulfur in wet
        desulfurization of gases)
IT
     7704-34-9P, Sulfur, preparation
     RL: BYP (Byproduct); PUR (Purification or recovery); PREP (Preparation)
        (method for recovering sulfur in wet desulfurization of gases)
ΙT
     497-19-8, Sodium carbonate, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (absorption solution component; in recovery of sulfur in wet
        desulfurization of gases)
RN
     497-19-8 HCA
CN
     Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)
```

О || НО— С— ОН

●2 Na

RN 7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

L90 ANSWER 17 OF 54 HCA COPYRIGHT 2004 ACS on STN

122:294669 Recovery of technical sulfur from concentrates resulted from flotation of ores containing sulfur. Fodor, Iosif; Puia, Mircea Laurean; Dajbukat, Martin Vasile (Combinatul Minier, Gura Humorului, Rom.). Rom. RO 103905 Bl 19940620, 5 pp. (Romanian). CODEN: RUXXA3. APPLICATION: RO 1989-139204 19890412.

AB To optimize the process and increase S yield, S is melted from the concs. containing 45-70% S by heating with live steam under known conditions in a an autoclave without stirring in the presence of ≥0.1% additive (relative to the solid content of the concentrate suspension) comprising preferably a mixture of Na tripolyphosphate and Na2CO3 in a 1.4:1 weight ratio and kerosene for pH control, at a ratio of total solids added to liquid 1.2:1.1. The process can be used independent of the source or nature of the starting ore.

IC ICM C01B017-033 ICS C01B017-05

CC 49-1 (Industrial Inorganic Chemicals)

ST sulfur recovery ore flotation conc; sodium tripolyphosphate
sulfur melt recovery; sodium carbonate sulfur
melt recovery; kerosene sulfur melt recovery

IT Kerosine

RL: TEM (Technical or engineered material use); USES (Uses) (pH regulator; recovery of tech. sulfur from concs. resulted from flotation of ores containing sulfur)

IT Recycling

(recovery of tech. sulfur from concs. resulted from flotation of ores containing sulfur)

IT Ore treatment

(flotation, concentrate; recovery of tech. sulfur from concs. resulted from flotation of ores containing sulfur)

IT 497-19-8, Sodium carbonate, uses 7758-29-4, Sodium tripolyphosphate

RL: TEM (Technical or engineered material use); USES (Uses) (pH regulator; recovery of tech. sulfur from concs. resulted from flotation of ores containing sulfur)

IT 7704-34-9P, Sulfur, preparation

RL: PUR (Purification or recovery); PREP (Preparation) (recovery of tech. sulfur from concs. resulted from flotation

of ores containing sulfur) IT 497-19-8, Sodium carbonate, uses RL: TEM (Technical or engineered material use); USES (Uses) (pH regulator; recovery of tech. sulfur from concs. resulted from flotation of ores containing sulfur)

RN497-19-8 HCA

Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME) CN

0 HO-C-OH

●2 Na

IT 7704-34-9P, Sulfur, preparation RL: PUR (Purification or recovery); PREP (Preparation) (recovery of tech. sulfur from concs. resulted from flotation of ores containing sulfur)

7704-34-9 HCA RN

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

L90 ANSWER 18 OF 54 HCA COPYRIGHT 2004 ACS on STN 122:191588 Flue Gas Desulfurization Scheme To Recover Elemental Sulfur. Sell, Nancy J.; Norman, Jack C.; Ciriacks, John A. (Department of Natural and Applied Sciences, University of Wisconsin, Green Bay, WI, 54311-7001, USA). Industrial & Engineering Chemistry Research, 34(4), 1428-33 (English) 1995. CODEN: IECRED. ISSN: 0888-5885. Publisher: American Chemical Society.

AΒ A sodium-based flue gas desulfurization process, employing reductive burning and a Claus recovery system, has the advantage of producing elemental sulfur as a byproduct. This study optimized the process conditions for two steps in the proposed reaction scheme: (1) removing approx. one-sixth of the sulfur in the spent scrubbing liquor by a . combination of acidification and evaporation and (2) precipitation of 50% of the

remaining Na2SO3 (s) by the addition of NaHCO3/Na2CO3 slurry. Several combinations of process conditions can be used to produce favorable results, but the optimum from a tech. perspective is to add concentrated H2SO4 to the spent liquor to 1.5% volume/volume, concentrate the mixture to 70% by weight, and

then treat it with a >70% solids slurry of NaHCO3/Na2CO3. The optimum from an economic perspective is dependent on the relative costs of H2SO4 and the steam needed to concentrate the spent liquor.

49-1 (Industrial Inorganic Chemicals) CC Section cross-reference(s): 59

ST flue gas desulfurization sulfur recovery

ΙT Flue gases

> (sodium-based flue gas desulfurization process to recover sulfur)

IT 144-55-8, Sodium bicarbonate, uses 497-19-8, Sodium carbonate, uses 7664-93-9, Sulfuric acid, uses

```
Hertzog
                                     10/688,243
     RL: NUU (Other use, unclassified); USES (Uses)
        (sodium-based flue gas desulfurization process to recover
        sulfur)
IT
     7704-34-9P, Sulfur, preparation
     RL: PUR (Purification or recovery); PREP (Preparation)
        (sodium-based flue gas desulfurization process to recover
        sulfur)
     144-55-8, Sodium bicarbonate, uses 497-19-8, Sodium
IT
     carbonate, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (sodium-based flue gas desulfurization process to recover
RN
     144-55-8 HCA
     Carbonic acid monosodium salt (8CI, 9CI) (CA INDEX NAME)
   0
HO- C- OH
 Na
RN
     497-19-8 HCA
CN
     Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)
```

•2 Na

HO-C-OH

S

```
L90 ANSWER 19 OF 54 HCA COPYRIGHT 2004 ACS on STN

121:86868 Process for recovering sulfur from carbon
dioxide- and hydrogen sulfide-containing acid gas mixtures. Towler, Gavin
P.; Lynn, Scott (University of California, USA). PCT Int. Appl. WO
9413579 A1 19940623, 18 pp. DESIGNATED STATES: W: AT, AU, BB,
BG, BR, BY, CA, CH, CZ, DE, DK, ES, FI, GB, HU, JP, KP, KR, KZ, LK, LU,
LV, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SK, UA, VN; RW: AT,
BE, BF, BJ, CF, CG, CH, CI, CM, DE, DK, ES, FR, GA, GB, GR, IE, IT, LU,
MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2.
APPLICATION: WO 1993-US11927 19931208. PRIORITY: US 1992-991163 19921216.

AB The process com prises contacting the the gas at ≥700° with
```

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a catalyst that enhances the thermal dissociation of H2S to H2 and S2. The
     equilibrium of the thermal decomposition of H2S is shifted by the
equilibration of
     the water-gas-shift reaction so as to favor the formation of elemental S.
     The primary products of the overall reaction are S2, CO, H2 and H2O.
     Small amts. of COS, SO2 and CS2 may also be formed. Rapid quenching of
     the reaction mixture to ≤600° results in a substantial
     increase in the efficiency of the conversion of H2S to S. Plant economy
     is further advanced by treating the product gases to remove byproduct COS
     by hydrolysis, whereby the COS is converted back to CO2 and H2S.
     ICM C01B017-04
ICS C01B017-16; C01B031-20; C01B031-26
IC
     49-1 (Industrial Inorganic Chemicals)
     Section cross-reference(s): 51
     Natural gas
IT
     RL: USES (Uses)
        (carbon dioxide- and hydrogen sulfide-containing, sulfur
        recovery from)
IT
     Transition metal sulfides
     RL: CAT (Catalyst use); USES (Uses)
        (decomposition catalysts, for sulfur recovery from
        carbon dioxide- and hydrogen sulfide-containing acid gas mixts.)
IT
     Alcohols, properties
     RL: PRP (Properties)
        (C1-4, amino, solns. containing potassium carbonate and,
        absorption in, of hydrogen sulfidfe, in sulfur
        recovery from carbon dioxide- and hydrogen sulfide-containing acid
        gas mixts.)
IT
     Alcohols, properties
     RL: PRP (Properties)
        (C1-4, iminodi-, solns. containing potassium carbonate and,
        absorption in, of hydrogen sulfide, in sulfur
        recovery from carbon dioxide- and hydrogen sulfide-containing acid
        gas mixts.)
IT
     1315-03-3, Vanadium sulfide (V2S3) 1317-33-5, Molybdenum disulfide, uses
     12018-22-3, Chromium sulfide (Cr2S3) 12138-09-9, Tungsten disulfide
     RL: CAT (Catalyst use); USES (Uses)
        (decomposition catalyst, for sulfur recovery from carbon
        dioxide- and hydrogen sulfide-containing acid gas mixts.)
     7783-06-4, Hydrogen sulfide, reactions
IT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (decomposition of, in presence of carbon dioxide, into hydrogen and
        sulfur, in natural gas purification)
IT
     7704-34-9P, Sulfur, preparation
     RL: PREP (Preparation)
        (formation of hydrogen and, by hydrogen sulfide decomposition in natural gas
        purification)
IT
     124-38-9, Carbon dioxide, uses
     RL: USES (Uses)
        (gas mixts. containing hydrogen sulfide and, sulfur
        recovery from)
IT
     463-58-1, Carbonyl sulfide
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (hydrolysis of, for carbon dioxide and hydrogen sulfide, in
        sulfur recovery from carbon dioxide- and hydrogen
        sulfide-containing acid gas mixts.)
     584-08-7P, Potassium carbonate
     RL: PREP (Preparation)
```

```
(solns. containing alkanolamines and dialkanolamines and, absorption in, of
        hydrogen sulfidfe, in sulfur recovery from carbon
        dioxide- and hydrogen sulfide-containing acid gas mixts.)
     7704-34-9P, Sulfur, preparation
     RL: PREP (Preparation)
        (formation of hydrogen and, by hydrogen sulfide decomposition in natural gas
        purification)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
TT
     124-38-9, Carbon dioxide, uses
     RL: USES (Uses)
        (gas mixts. containing hydrogen sulfide and, sulfur
        recovery from)
RN
     124-38-9 HCA
CN
     Carbon dioxide (8CI, 9CI) (CA INDEX NAME)
o = c = o
     584-08-7P, Potassium carbonate
     RL: PREP (Preparation)
        (solns. containing alkanolamines and dialkanolamines and, absorption in, of
        hydrogen sulfidfe, in sulfur recovery from carbon
        dioxide- and hydrogen sulfide-containing acid gas mixts.)
RN
     584-08-7 HCA
CN
     Carbonic acid, dipotassium salt (8CI, 9CI) (CA INDEX NAME)
   O
HO-C-OH
 ●2 K
L90 ANSWER 20 OF 54 HCA COPYRIGHT 2004 ACS on STN
119:274498 Purification of sulfur. Ma, Yongming; Han,
     Yuying; Yan, Yixin (Shanxi University, Peop. Rep. China). Faming Zhuanli
     Shenqing Gongkai Shuomingshu CN 1073657 A 19930630, 5 pp.
```

(Chinese). CODEN: CNXXEV. APPLICATION: CN 1991-111804 19911222. The process comprises placing the S-containing material in a vessel heated at AB 115-445°, introducing water or steam into the vessel to form a gas

mixture containing S and water, cooling the gas mixture to precipitate S, and

washing and

drying the S. Additive corresponding to 0.01-10% of the S content in the S-containing material may be added to the system for increased efficiency, and the additive is NH3, urea or NH4 salt of a nonoxidizing acid selected from ≥1 of (NH4)2CO3, NH4HCO3, NH4C1, NH4H2PO4, (NH4)2HPO4. The S-containing material is industrial S with high As content, or recovered crude S from industries. The steam is may be superheated steam.

IC ICM C01B017-02

49-1 (Industrial Inorganic Chemicals)

```
ST
     sulfur purifn
IT
     Steam
        (purification with, of crude sulfur, additives for, for
        increased efficiency)
     57-13-6, Urea, uses 506-87-6, Ammonium carbonate
IT
     1066-33-7, Ammonium bicarbonate 7664-41-7, Ammonia, uses
     7722-76-1 7783-28-0, Ammonium phosphate, dibasic 12125-02-9, Ammonium
     chloride, uses
     RL: USES (Uses)
        (additive, in sulfur purification by steam-treating, for
        increased efficiency)
ΙT
     7704-34-9P, Sulfur, preparation
     RL: PREP (Preparation)
        (purification of crude, by steam-treating, additives for increased
        efficiency in)
ΙT
     7440-38-2, Arsenic, miscellaneous
     RL: REM (Removal or disposal); PROC (Process)
        (removal of, from industrial sulfur, in sulfur
        purification)
ΙT
     7732-18-5
     RL: USES (Uses)
        (steam, purification with, of crude sulfur, additives
        for, for increased efficiency)
IT
     506-87-6, Ammonium carbonate 1066-33-7,
     Ammonium bicarbonate
     RL: USES (Uses)
        (additive, in sulfur purification by steam-treating, for
        increased efficiency)
     506-87-6 HCA
RN
     Carbonic acid, diammonium salt (8CI, 9CI) (CA INDEX NAME)
HO- C- OH
●2 NH3
     1066-33-7 HCA
     Carbonic acid, monoammonium salt (8CI, 9CI) (CA INDEX NAME)
HO- C- OH
 ● инз
IT
     7704-34-9P, Sulfur, preparation
     RL: PREP (Preparation)
        (purification of crude, by steam-treating, additives for increased
        efficiency in)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI)
                       (CA INDEX NAME)
```

L90 ANSWER 22 OF 54 HCA COPYRIGHT 2004 ACS on STN

S

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116:27118 Removal and recovery of sulfur dioxide from
     waste gases, especially from metallurgical plants, and from flue gases.
     Matschiner, Hermann; Kain, Christoph; Haase, Rudolf; Maschmeier, Claus
     Peter (Martin-Luther-Universitaet Halle-Wittenberg, Germany). Ger. (East)
     DD 293999 A5 19910919, 4 pp. (German). CODEN: GEXXA8.
     APPLICATION: DD 1989-334815 19891124.
AΒ
     SO2 is removed from flue or waste gases by scrubbing with a sec. amine,
     e.g., dimethylamine, at 30-80°. The SO2-amine adducts are used to
     manufacture sulfites or bisulfites by reaction with aqueous alkali hydroxides,
     carbonates, or bicarbonates. Addnl. S may be used to manufacture
     thiosulfates. The adduct may be reacted with H2S to form S. The amine is
     recovered for recycling.
     ICM C01B017-60
     ICS C01B017-62; C01B017-98; C01B017-64; B01D053-34
     59-4 (Air Pollution and Industrial Hygiene)
     Section cross-reference(s): 49, 55, 56
     sulfur dioxide recovery flue gas; sulfite manuf sulfur
     dioxide waste; bisulfite manuf sulfur dioxide waste; thiosulfate manuf
     sulfur dioxide waste; sulfur manuf sulfur dioxide waste
IT
     1310-73-2, Sodium hydroxide, uses
     RL: USES (Uses)
        (in recovery of sulfur compds. from waste gases
        containing sulfur dioxide, dimethylamine in)
ΙT
     7783-06-4, Hydrogen sulfide, uses
     RL: USES (Uses)
        (in recovery of sulfur from waste gases containing
        sulfur dioxide, dimethylamine in)
ΙT
     124-40-3, Dimethylamine, uses
     RL: USES (Uses)
        (in removal and recovery of sulfur dioxide from
        waste gases)
ΙT
     463-79-6DP, Carbonic acid, alkali salts 7704-34-9P,
     Sulfur, preparation 7772-98-7P, Sodium thiosulfate
     7782-99-2DP, Sulfurous acid, alkali metal salts 13686-28-7DP,
     Thiosulfuric acid, alkali metal salts
     RL: PREP (Preparation)
        (recovery of, from waste gases containing sulfur dioxide,
        dimethylamine in)
IT
     463-79-6DP, Carbonic acid, alkali salts 7704-34-9P,
     Sulfur, preparation
     RL: PREP (Preparation)
        (recovery of, from waste gases containing sulfur dioxide,
        dimethylamine in)
RM
     463-79-6 HCA
     Carbonic acid (7CI, 8CI, 9CI) (CA INDEX NAME)
CN
HO-C-OH
```

RN

7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

L90 ANSWER 24 OF 54 HCA COPYRIGHT 2004 ACS on STN
113:117877 Manufacture of sublimed amorphous sulfur. Krajewski,
Jan; Gatarz, Zdzislawa; Dojka, Marian; Krawczyk, Boguslaw; Pantula,
Zbigniew; Warzybok, Jozef; Malarz, Janusz; Kopec, Jozef (Osrodek
Badawczo-Rozwojowy Przemyslu Siarkowego "Siarkopol", Pol.). Pol. PL
148229 Bl 19890930, 5 pp. Abstracted and indexed from the
unexamined application. (Polish). CODEN: POXXA7. APPLICATION: PL
1987-268208 19871013.

AB S-containing air from S mines is introduced into absorption column and S vapor is contacted with counter current-sprayed **liquid**, preferably aqueous inorg. salts with alkaline reaction, and the resulting S suspension is separated

by known methods, washed, filtered and dried to obtain fine S powder having globular shape.

IC ICM C01B017-10

CC 49-1 (Industrial Inorganic Chemicals)

ST amorphous sulfur manuf

IT 7704-34-9P, Sulfur, preparation

RL: PREP (Preparation)

(manufacture of amorphous)

IT 497-19-8, Sodium carbonate, uses and miscellaneous RL: USES (Uses)

(sprayable liquid containing, in sulfur manufacture by resublimation)

IT 7704-34-9P, Sulfur, preparation

RL: PREP (Preparation)

(manufacture of amorphous)

RN 7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

RN 497-19-8 HCA

CN Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)

но— с— он || О

●2 Na

L90 ANSWER 26 OF 54 HCA COPYRIGHT 2004 ACS on STN 112:215619 Production of water-suspendable liquid or solid

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sulfur concentrates. Teren, Jan; Nadvornik, Robert; Hutar,
     Eduard; Stanik, Vojtech (Czech.). Czech. CS 250858 Bl 19880815,
     8 pp. (Slovak). CODEN: CZXXA9. APPLICATION: CS 1983-5514 19830725.
     Molten S is emulsified in sulfite waste liquor and/or a liquosulfonic acid
     salt containing \geq 1 Ca2+, Mg2+, Fe2+, Fe3+, Cu2+, Mn2+, Zn2+, Co2+, and
     TiO2+. Optionally, the sulfite waste liquor is preheated to
     65-105°. The S concentrate is a fertilizer. A concentrate was prepared by
     mixing concentrated sulfite waste liquor (dry matter 52.7%), 546.2, MgSO4.7H2O 93.9, and H2O 72.1 g, at 85°. After settling, the solution (499.7 g)
     was preheated to 95-105°, and molten S 283.1 g was added. After
     drying, the solid concentrate contained total S 56.34, total Mg 1.41,
water-soluble
     Mg 1.40, and total Ca 0.30 weight%.
     ICM C01B017-00
ΙC
     19-6 (Fertilizers, Soils, and Plant Nutrition)
CC
     7704-34-9P, Sulfur, biological studies
IT
     RL: AGR (Agricultural use); BIOL (Biological study); PREP (Preparation);
     USES (Uses)
        (fertilizer, manufacture of, by emulsification of melt, in sulfite waste
        liquid and/or lignosulfonate)
     546-46-3, Zinc citrate 546-93-0, Magnesium
IT
     carbonate (MqCO3)
                         1309-48-4, Magnesia (MgO), biological studies
     7439-89-6D, Iron, salts 7439-96-5D, Manganese, salts 7440-32-6D,
     Titanium, salts 7440-48-4D, Cobalt, salts
                                                   7440-50-8D, Copper, salts
     7440-66-6D, Zinc, salts 7487-88-9, Sulfuric acid magnesium salt (1:1),
     biological studies 7720-78-7 7733-02-0, Zinc sulfate (ZnSO4)
     7758-98-7, Sulfuric acid copper(2+) salt (1:1), biological studies
     8062-15-5D, Lignosulfonic acid, salts 12519-36-7, Zinc EDTA 15498-89-2
     RL: BIOL (Biological study)
        (in sulfur fertilizer concentrate manufacture)
IΤ
     7704-34-9P, Sulfur, biological studies
     RL: AGR (Agricultural use); BIOL (Biological study); PREP (Preparation);
     USES (Uses)
        (fertilizer, manufacture of, by emulsification of melt, in sulfite waste
        liquid and/or lignosulfonate)
     7704-34-9 HCA
RN
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
     546-46-3, Zinc citrate 546-93-0, Magnesium
     carbonate (MgCO3)
     RL: BIOL (Biological study)
        (in sulfur fertilizer concentrate manufacture)
RN
CN
     1,2,3-Propanetricarboxylic acid, 2-hydroxy-, zinc salt (2:3) (9CI) (CA
     INDEX NAME)
```

$$\begin{array}{c} \text{CO}_2\text{H} \\ | \\ \text{HO}_2\text{C} - \text{CH}_2 - \text{C} - \text{CH}_2 - \text{CO}_2\text{H} \\ | \\ \text{OH} \end{array}$$

●3/2 Zn

RN 546-93-0 HCA

CN Carbonic acid, magnesium salt (1:1) (8CI, 9CI) (CA INDEX NAME)

O . || HO— C— OH

Mq

L90 ANSWER 27 OF 54 HCA COPYRIGHT 2004 ACS on STN

112:23522 MHD seed recovery and regeneration. McIlroy, R. A.; Probert, P. B.; Jackson, D. M.; Lahoda, E. J. (Babcock and Wilcox Co., USA). Proceedings of the Intersociety Energy Conversion Engineering Conference, 24th(Vol. 2), 1025-31 (English) 1989. CODEN: PIECDE. ISSN: 0146-955X.

AB A process is described for regeneration of K2CO3 seed from spent material recovered as K2SO4 from the residues of coal-fired MHD generators; the process is based on reduction of SO42- followed by separation of S and K.

Reduction is

carried out in a rotary kiln by combustion of K2SO4-coal mixts. (2:1) at 700°; K2S2 forms and is dissolved to form a green liquor which is clarified and then carbonated to H2S and K2CO3, by a modified Tampella process. The H2S is converted to elemental S using the Claus process; K2CO3 is purified by crystallization to 95% purity, with Na and Cl as main impurities. The flue gas from the kiln is treated in scrubbers to recover H2S; the plant also has waste heat recovery systems to improve the energy balance.

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): **49**, 51, 59

potassium carbonate seed MHD recovery; MHD generator potassium seed recovery; redn potassium sulfate coal kiln; carbonation potassium MHD seed recovery; flue gas potassium sulfate redn; waste heat MHD seed recovery

IT Air pollution

(by flue gases, from potassium sulfate-coal combustion, in potassium carbonate MHD seed recovery process, abatement of)

IT Flue gases

(from potassium sulfate-coal combustion, in potassium carbonate MHD seed recovery process, cleaning of)

IT Ashes (residues)

(from potassium sulfate-coal combustion, in potassium carbonate MHD seed recovery process, disposal of)

IT Reduction

```
(of potassium sulfate, by coal, in rotary kiln, for potassium
        carbonate MHD generator seed recovery)
IT
     Electric generators
         (MHD, potassium carbonate seed for, recovery of,
        reduction/carbonation process for)
IT
         (waste, recovery of, from potassium sulfate-coal combustion, in
        potassium carbonate MHD seed recovery process)
IT
     7783-06-4P, Hydrogen sulfide, preparation
     RL: RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)
         (formation and decomposition of, sulfur recovery by, in
        recovery/regeneration of potassium carbonate, for MHD
        generator seed)
     7704-34-9P, Sulfur, preparation
TT
     RL: FORM (Formation, nonpreparative); PREP (Preparation)
        (formation of, in recovery of potassium carbonate, for MHD
        generator seed)
IT
     584-08-7P, Potassium carbonate (K2CO3)
     RL: PREP (Preparation)
        (recovery of, potassium sulfate reduction/carbonation process for, for MHD
        generator seed)
IT
     7778-80-5, Potassium sulfate (K2SO4), reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reduction of, by coal, in rotary kiln, for potassium carbonate
        MHD generator seed recovery)
IT
     7704-34-9P, Sulfur, preparation
     RL: FORM (Formation, nonpreparative); PREP (Preparation)
        (formation of, in recovery of potassium carbonate, for MHD
        generator seed)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
     584-08-7P, Potassium carbonate (K2CO3)
IT
     RL: PREP (Preparation)
        (recovery of, potassium sulfate reduction/carbonation process for, for MHD
        generator seed)
RN
     584-08-7 HCA
     Carbonic acid, dipotassium salt (8CI, 9CI) (CA INDEX NAME)
HO-C-OH
 ●2 K
L90 ANSWER 28 OF 54 HCA COPYRIGHT 2004 ACS on STN
111:197844 Process for removal of hydrogen sulfide from sour gas streams.
     Byers, Dallas L. (Shell Oil Co., USA). U.S. US 4840782 A
     19890620, 9 pp. Cont. of U.S. Ser. No. 937,446, abandoned.
```

PRIORITY: US 1984-614363 19840529; US 1985-790763 19851023; US 1986-937446

(English). CODEN: USXXAM. APPLICATION: US 1987-139140 19871221.

19861205.

AB H2S is removed from sour gas by contacting the sour gas stream in a contacting zone with an aqueous reaction solution substantially free of anthraquinone disulfonate (<0.2 g/L) and containing an effective amount of V5+ ions to oxidize H2S, producing a sweet gas stream and an aqueous solution containing

solid S and V4+ ions. The reaction solution has a pH of 8-11 and also contains an amount of phosphate ions sufficient to provide a molar ratio of phosphate ions to V4+ ions ≥0.1. The solid S is removed from the aqueous solution and the remaining aqueous solution is regenerated and recycled.

IC ICM C01B017-05

NCL 423576600

CC 48-1 (Unit Operations and Processes)

Section cross-reference(s): 49, 51

IT Fuel gases

Natural gas

RL: USES (Uses)

(hydrogen sulfide removal from, by oxidation with solution containing vanadium ions and phosphate ions)

107-70-0 108-10-1, Methyl isobutyl ketone 108-32-7, Propylene carbonate 108-94-1, Cyclohexanone, uses and miscellaneous 110-13-4, 2,5-Hexanedione 111-46-6, Diethylene glycol, uses and miscellaneous 111-90-0, Diethylene glycol monoethyl ether 112-60-7, Tetraethylene glycol 123-42-2, Diacetone alcohol 123-54-6, 2,4-Pentanedione, uses and miscellaneous 126-33-0, Sulfolane 141-79-7, Mesityl oxide 142-92-7, Hexyl acetate 143-24-8, Tetraethylene glycoldimethyl ether 872-50-4, N-Methyl pyrrolidone, uses and miscellaneous

RL: USES (Uses)

(absorbent, for removal of hydrogen sulfide from sour gas streams)

IT 7704-34-9P, Sulfur, preparation

RL: FORM (Formation, nonpreparative); PREP (Preparation) (formation of, from hydrogen sulfide oxidation with

vanadium ion containing solution, in sour gas treatment)

TT 7632-05-5, Sodium phosphate 10124-31-9, Ammonium phosphate 10377-52-3, Lithium phosphate 11105-06-9, Sodium vanadate 11115-67-6, Ammonium vanadate 11126-15-1, Lithium vanadium oxide 11126-20-8, Potassium vanadate 16068-46-5, Potassium phosphate

RL: USES (Uses)

(in removal of hydrogen **sulfide** by **oxidation**, from sour gas streams)

IT 124-38-9, Carbon dioxide, uses and miscellaneous

RL: REM (Removal or disposal); PROC (Process)

(removal of, from sour gas containing hydrogen sulfide)

IT 108-32-7, Propylene carbonate 110-13-4,

2,5-Hexanedione 123-54-6, 2,4-Pentanedione, uses and

miscellaneous RL: USES (Uses)

(absorbent, for removal of hydrogen sulfide from sour gas streams)

RN 108-32-7 HCA

CN 1,3-Dioxolan-2-one, 4-methyl- (9CI) (CA INDEX NAME)

RN 110-13-4 HCA

CN 2,5-Hexanedione (8CI, 9CI) (CA INDEX NAME)

RN 123-54-6 HCA

CN 2,4-Pentanedione (8CI, 9CI) (CA INDEX NAME)

$$\begin{array}{c|c} \mathsf{O} & \mathsf{O} \\ \parallel & \parallel \\ \mathsf{Me}\mathsf{-C}\mathsf{-CH}_2\mathsf{-C}\mathsf{-Me} \end{array}$$

IT 7704-34-9P, Sulfur, preparation

RL: FORM (Formation, nonpreparative); PREP (Preparation) (formation of, from hydrogen sulfide oxidation with vanadium ion containing solution, in sour gas treatment)

RN 7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

RN. 124-38-9 HCA

CN Carbon dioxide (8CI, 9CI) (CA INDEX NAME)

o== c== o

L90 ANSWER 30 OF 54 HCA COPYRIGHT 2004 ACS on STN
109:112575 Effect of the method of copper oxide addition on chemisorption
 properties of zinc-containing sulfur-purification
 materials. Dantsig, G. A.; Grechenko, A. N.; Grigorev, V. V.; Serova, L.
 P.; Yagodkina, G. N. (Gos. Inst. Azotn. Prom. Prod. Org. Sint.,
 Novomoskovsk, USSR). Zhurnal Prikladnoi Khimii (Sankt-Peterburg, Russian
 Federation), 61(6), 1240-6 (Russian) 1988. CODEN: ZPKHAB.
 ISSN: 0044-4618.

 $\ensuremath{\mathsf{AB}}$ $\ensuremath{\mathsf{The}}$ chemisorption of H2S and other S-containing materials on S-purification agents

(oxides, e.g., GIAP-10-2) occurs 50-70° lower than on activated ZnO. The effect of CuO on the sorption properties of S-purification agents was studied using a mixture of basic carbonates of Zn and Cu and a mixture of ZnO and CuO treated with (NH4)2CO3. The presence of Cu in the mixture significantly enhanced the decomposition of H2S before chemisorption. The (NH4)2CO3 treatment of CuO and ZnO resulted in the formation of double salts, which, after being reduced by H2, facilitated the introduction of Cu into the ZnO lattice. The Cu in the S-purification agents acts as a promoter, enhancing the reaction between H2S and ZnO.

CC 48-1 (Unit Operations and Processes)
 Section cross-reference(s): 49

ST copper oxide hydrogen sulfide sorbent; zinc oxide hydrogen sulfide

```
sorbent; sulfur purifn material hydrogen
sulfide sorbent

IT 1317-38-0, Copper oxide, uses and miscellaneous
RL: USES (Uses)
```

(chemisorption of hydrogen sulfide on zinc-containing sulfurpurification agents in relation to)

IT 7783-06-4, Hydrogen sulfide, reactions
RL: RCT (Reactant); RACT (Reactant or reagent)

(chemisorption of, on zinc-containing sulfur-purification agents, copper oxide effect on)

IT 506-87-6, Ammonium carbonate

RL: USES (Uses)

(in preparation of zinc oxide, for purification of sulfur)

IT 7704-34-9P, Sulfur, preparation

RL: PUR (Purification or recovery); PREP (Preparation)
 (purification of, zinc-containing agent for, hydrogen sulfide chemisorption by)

IT 506-87-6, Ammonium carbonate

RL: USES (Uses)

(in preparation of zinc oxide, for purification of sulfur)

RN 506-87-6 HCA

CN Carbonic acid, diammonium salt (8CI, 9CI) (CA INDEX NAME)

●2 NH3

RN 7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

L90 ANSWER 31 OF 54 HCA COPYRIGHT 2004 ACS on STN
109:60724 Process for removing hydrogen sulfide from gas streams. De Haan,
Robert; Dry, Mark Eberhard; Dressler, Flemming Hermann; Hesse, Horst
Joachim Franz August (Sasol Operations (Pty.) Ltd., S. Afr.). S. African
ZA 8703112 A 19871230, 25 pp. (English). CODEN: SFXXAB.
APPLICATION: ZA 1987-3112 19870430. PRIORITY: ZA 1986-3264 19860501.
AB H2S is removed from waste gases by scrubbing with an alkaline solution
containing

oxovanadium (V), with alkalinity present as OH-, CO32-, and HCO3- to maintain the pH at 7.4-9 where CO32- in <12.5 g/L. The oxovanadium (IV) produced from S oxidation can be oxidized to the pentavalent form by bubbling an O-containing gas through the scrubbing solution A scrubbing solution containing

pentavalent V 1.9, Na2CO3 40, diethanol amine 40, and NaSCN 20 g/L was used to remove 97.7% of H2O present at 1% in CO2, for a solution loading of

300 mg H2S/L. The S formed was recovered from froth in the oxidizer.

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IC
     ICM C01B
     ICS B01D; C07C; C01G
CC
     59-4 (Air Pollution and Industrial Hygiene)
     Section cross-reference(s): 49
     hydrogen sulfide scrubbing waste gas; sulfur recovery
ST
     hydrogen sulfide scrubbing; vanadium pentoxide hydrogen sulfide
     scrubbing
TT
     16408-26-7
     RL: OCCU (Occurrence)
         (hydrogen sulfide removal from gas by scrubbing with, with
        sulfur recovery)
IT
     7704-34-9P, Sulfur, preparation
     RL: PREP (Preparation)
         (recovery of, in hydrogen sulfide scrubbing from
        gas with vanadium pentoxide)
IT
     7783-06-4, Hydrogen sulfide, uses and miscellaneous
     RL: REM (Removal or disposal); PROC (Process)
        (removal of, from gas, scrubbing with vanadium pentoxide for,
        sulfur recovery in)
IT
     111-42-2, Diethanol amine, uses and miscellaneous 497-19-8,
     Sodium carbonate, uses and miscellaneous
                                                 540-72-7, Sodium
     thiocyanate
     RL: USES (Uses)
         (scrubbing solution containing, for hydrogen sulfide removal from gas)
ΙT
     7704-34-9P, Sulfur, preparation
     RL: PREP (Preparation)
        (recovery of, in hydrogen sulfide scrubbing from
        gas with vanadium pentoxide)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
ΙT
     497-19-8, Sodium carbonate, uses and miscellaneous
     RL: USES (Uses)
        (scrubbing solution containing, for hydrogen sulfide removal from gas)
RN
     497-19-8 HCA
     Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)
   0
HO- C- OH
 ●2 Na
L90 ANSWER 33 OF 54 HCA COPYRIGHT 2004 ACS on STN
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John Calve EIC- 1700

AΒ

source and pyritic concentrates, in dry form at atmospheric pressure. Saddy, Maury (Centro de Tecnologia Promon-CTP, Brazil). Braz. Pedido PI BR 8506126 A 19870630, 6 pp. (Portuguese). CODEN: BPXXDX.

The title process includes thermal decomposition of pyrite in a

107:239266 Preparation of elementary sulfur from pyrites of any

APPLICATION: BR 1985-6126 19851206.

```
{f fluidized} bed at 400-1200° to produce S by condensation of
     S vapor as well as a mixture of pyrrhotite and diverse residues, magnetic
     separation of the pyrrhotite from the other residues for recycling as a feed to
     the fluidized bed, operating with an excess of air at
     400-1200° to produce Fe2O3 and SO2. Fuel is fed to a combustion
     reactor in the fluidized bed, operating with 20% excess air at
     600-1200°, and 100% excess CaCO3 is added to react with the toxic
     SO2 to produce CaSO4, CaO, and CO2. The heat of the gases from combustion
     and roasting may be partially used in the 1st and 2nd stages of the
     process for drying and thermal decomposition
     ICM C22B001-10
     ICS C01B017-00
     49-1 (Industrial Inorganic Chemicals)
     sulfur prodn pyrite thermal decompn; ferric oxide byproduct
     sulfur prodn; calcium sulfate byproduct sulfur prodn;
     calcium oxide byproduct sulfur prodn; carbon dioxide formation
     sulfur prodn
     1310-50-5P, Pyrrhotite
     RL: RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)
        (formation and thermal decomposition of, in sulfur production from
        pyrite)
     1309-37-1P, Ferric oxide, preparation
                                             7446-09-5P, Sulfur
     dioxide, preparation
     RL: FORM (Formation, nonpreparative); PREP (Preparation)
        (formation of, in thermal decomposition of pyrrhotite, in sulfur
        production from pyrite)
     124-38-9P, Carbon dioxide, preparation 1305-78-8P, Calcium
                          7778-18-9P, Calcium sulfate
     oxide, preparation
     RL: PREP (Preparation)
        (formation, of, by calcium carbonate reaction with
        sulfur dioxide from pyrrhotite thermal decomposition in
        sulfur production from pyrite)
     7704-34-9P, Sulfur, preparation
     RL: PREP (Preparation)
        (production of, from pyrite, by thermal decomposition)
     471-34-1P, Calcium carbonate, reactions
     RL: RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)
        (reaction of, with sulfur dioxide from pyrrhotite thermal
        decomposition in sulfur production from pyrite)
     1309-36-0P, Pyrite, reactions
     RL: RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)
        (thermal decomposition of, sulfur production and phyrrhotite formation
     124-38-9P, Carbon dioxide, preparation
     RL: PREP (Preparation)
        (formation, of, by calcium carbonate reaction with
        sulfur dioxide from pyrrhotite thermal decomposition in
        sulfur production from pyrite)
     124-38-9 HCA
     Carbon dioxide (8CI, 9CI) (CA INDEX NAME)
o = c = 0
```

CC

ST

IΤ

IT

IT

IT

TΤ

IT

IT

RN

CN

IT 7704-34-9P, Sulfur, preparation RL: PREP (Preparation) (production of, from pyrite, by thermal decomposition) RN7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

RN 471-34-1 HCA

CN Carbonic acid calcium salt (1:1) (8CI, 9CI) (CA INDEX NAME)

HO-C-OH

Ca

L90 ANSWER 34 OF 54 HCA COPYRIGHT 2004 ACS on STN
107:83651 Method for purifying stratal and drainage water of sulfur
pits. Gorshkov, V. P.; Salyuk, A. P. (USSR). U.S.S.R. SU 1296517 A1
19870315 From: Otkrytiya, Izobret. 1987, (10), 104. (Russian).
CODEN: URXXAF. APPLICATION: SU 1985-3852230 19850206.

AB Title method includes removal of suspended and colloidal contaminants, air stripping of H2s with subsequent chemical treatment of the air-gas mixture and separation of the treated air, electrochem.

treatment of the water, anion exchange filtration, and regeneration of the anion exchanger. For a no-waste process, a decrease in reagent consumption, and use of the treated water as heat carrier for underground melting of S, after the suspended and colloidal contaminants are removed hydroacoustically as concentrated product and electrochem. treatment of clarified water in an electrolyzer, the anolyte is fed to the stripping stage and anion exchanged and the catholyte is mixed with the air-gas mixture after the chemical treatment stage and after removal of CaCOo and MgCOoin a sand filter, part of it is recombined with the treated anolyte and use for heat exchange. The other part of the treated catholyte is used to regenerate the anion exchanger and the spent regeneration solution is used to wash the sand filter, then containing CaCO4 and MgCO4, used for chemical treatment of the air-gas mixture, and mixed with the concentrated products centrifuged. The sediment is discharged and the centrifugate is mixed with incoming water. The sep. treated air, after catholyte mixing with the air-gas mixture, is recycled to the stripping stage.

IC ICM C02F009-00

CC 61-5 (Water)

Section cross-reference(s): 49

ST waste free **sulfur** pit water treatment; hydrogen **sulfide** air stripping water; electrochem treatment **sulfur** pit water; anion exchange **sulfur** pit water

IT Water purification

(anion exchange, in multistage treatment of **sulfur** pit stratal and drainage waters, with anion exchanger regeneration)

IT Water purification (electrodialysis, anolyte and catholyte from, recycling of, in

```
treatment of sulfur pit stratal and drainage waters)
ΙT
     Water purification
         (stripping, air, hydrogen sulfide removal in, of
        sulfur pit stratal and drainage water)
IT
     7704-34-9P, Sulfur, uses and miscellaneous
     RL: PREP (Preparation); USES (Uses)
         (drainage and stratal waters from pits for production of, multistage
        treatment of, waste free)
IT
     471-34-1, Calcium carbonate, uses and miscellaneous
     546-93-0, Magnesium carbonate
     RL: REM (Removal or disposal); PROC (Process)
        (removal of, from catholyte mixture with hydrogen sulfide
        -containing stripping air, in sulfur pit water treatment)
     7783-06-4, Hydrogen sulfide, uses and miscellaneous
IT
     RL: REM (Removal or disposal); PROC (Process)
        (removal of, from sulfur pit stratal and drainage water, air
        stripping for)
IT
     7704-34-9P, Sulfur, uses and miscellaneous
     RL: PREP (Preparation); USES (Uses)
        (drainage and stratal waters from pits for production of, multistage
        treatment of, waste free)
RN
     7704-34-9 HCA
     Sulfur (8CI, 9CI) (CA INDEX NAME)
CN
S
IT
     471-34-1, Calcium carbonate, uses and miscellaneous
     546-93-0, Magnesium carbonate
     RL: REM (Removal or disposal); PROC (Process)
        (removal of, from catholyte mixture with hydrogen sulfide
        -containing stripping air, in sulfur pit water treatment)
RN
     471-34-1 HCA
     Carbonic acid calcium salt (1:1) (8CI, 9CI) (CA INDEX NAME)
CN
HO-C-OH
  ● Ca
RN
     546-93-0 HCA
     Carbonic acid, magnesium salt (1:1) (8CI, 9CI) (CA INDEX NAME)
HO- C- OH
 Mg
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L90 ANSWER 36 OF 54 HCA COPYRIGHT 2004 ACS on STN

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106:122369 Method of winning of elemental sulfur from
     sulfur ore. Bylo, Zbigniew; Kolenda, Zygmunt; Jarosinska, Krystyna;
     Malinowska, Kazimiera; Krajewski, Jan; Krzeminski, Marian; Bold,
     Stanislaw; Kosinski, Zenobiusz (Akademia Gorniczo-Hutnicza, Pol.; Osrodek
     Badawczo-Rozwojowy Przemyslu Siarkowego "Siarkopol"). Pol. PL 127859 B1
     19860210, 2 pp. (Polish). CODEN: POXXA7. APPLICATION: PL
     1981-230270 19810319.
     After conventional flotation of a S ore (containing .apprx.18% S), the
     remelting S concentrate is leached to remove CaCO3. Leaching is done by
     SO2 for 1-3 h through a slurry containing 1-2.5 ton water/ton S concentrate at
     293-313 K. The S concentrate is separated from the formed Ca(HSO3)2 solution
     used for cellulose manufacture Then, the S concentrate in subjected to
conventional
     pressure-filtration refining. Thus, 200 kg S concentrate containing 76.45% S
     mixed with 500 kg water, and the resulting slurry was blown 60 min with
     SO2 at 293 K. The S concentrate was separated from the resulting Ca(HSO3)2
     washed, and dried at 378 K. The S content in the concentrate was 85.76%.
     C01B017-027
IC
CC
     49-1 (Industrial Inorganic Chemicals)
     Section cross-reference(s): 43
ST
     sulfur recovery ore refining; calcium
     carbonate removal sulfur conc; sulfite calcium hydrogen sulfur
ΤТ
     Leaching
        (of sulfur concentrate for calcium carbonate removal)
TΤ
     13780-03-5P
     RL: FORM (Formation, nonpreparative); PREP (Preparation)
        (formation of, in removal of calcium carbonate from sulfur
        concentrate by leaching with sulfur dioxide)
ΙT
     7446-09-5, Sulfur dioxide, uses and miscellaneous
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (leaching by, of sulfur concentrate, for calcium carbonate removal)
IT
     7704-34-9P, Sulfur, preparation
     RL: PREP (Preparation)
        (recovery of, from ore, concentrate leaching for calcium
        carbonate removal in relation to)
IT
     471-34-1, Calcium carbonate, uses and miscellaneous
     RL: REM (Removal or disposal); PROC (Process)
        (removal of, from sulfur concentrate, by leaching with sulfur dioxide)
IT
     7704-34-9P, Sulfur, preparation
     RL: PREP (Preparation)
        (recovery of, from ore, concentrate leaching for calcium
        carbonate removal in relation to)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
IT
     471-34-1, Calcium carbonate, uses and miscellaneous
     RL: REM (Removal or disposal); PROC (Process)
```

John Calve EIC- 1700

RNCN

Carbonic acid calcium salt (1:1) (8CI, 9CI) (CA INDEX NAME)

(removal of, from sulfur concentrate, by leaching with sulfur dioxide)

HO-C-OH

Ca

L90 ANSWER 37 OF 54 HCA COPYRIGHT 2004 ACS on STN 105:26587 Winning of pure sulfur from sulfur ore. Bylo, Zbigniew; Kolenda, Zygmunt; Jarosinska, Krystyna; Malinowska, Kazimiera; Krajewski, Jan; Krzeminski, Marian; Bold, Stanislaw; Kosinski, Zenobiusz (Akademia Gorniczo-Hutnicza, Pol.; Osrodek Badawczo-Rozwojowy Przemyslu Siarkowego "Siarkopol"). Pol. PL 127849 Bl 19831130, 2 pp. (Polish). CODEN: POXXA7. APPLICATION: PL 1981-230047 19810306. The yield of S was increased and the weight of carbonate residues was decreased when CO2 was passed through S concentrate obtained from an ore containing 18% elemental S. Thus, 200 kg S concentrate (76.45% elemental S) kg water were stirred at 293 K while CO2 was bubbled through for 60 min. Filtration, washing of filter cake, and drying gave enriched S concentrate containing 81.24% elemental S. ΙC C01B017-027 CC49-1 (Industrial Inorganic Chemicals) Section cross-reference(s): 54 STsulfur conc carbonate removal carbonation IT **7704-34-9P**, preparation RL: PREP (Preparation) (concentration of, from sulfur-containing ores, by carbon dioxide treatment) ΙT 124-38-9, uses and miscellaneous RL: USES (Uses) (sulfur-containing ores treated with, for sulfur enrichment) IT **7704-34-9P**, preparation RL: PREP (Preparation) (concentration of, from sulfur-containing ores, by carbon dioxide treatment) 7704-34-9 HCA RNSulfur (8CI, 9CI) (CA INDEX NAME) CN S TT 124-38-9, uses and miscellaneous RL: USES (Uses) (sulfur-containing ores treated with, for sulfur enrichment) RN 124-38-9 HCA Carbon dioxide (8CI, 9CI) (CA INDEX NAME) CN o = c = o

L90 ANSWER 38 OF 54 HCA COPYRIGHT 2004 ACS on STN 102:81133 Sulfur and calcium carbonate from gypsum.

Batista de Queiroz, Agnaldo; Peres, Luciano dos Santos (Fundacao Instituto Tecnologico do Estado de Pernambuco, Brazil). Braz. Pedido PI BR 8300949 A 19841002, 12 pp. (Portuguese). CODEN: BPXXDX. APPLICATION: BR 1983-949 19830113.

S and CaCO3 are produced from gypsum in a multistage process. Crushed AΒ gypsum is calcined to the hemihydrate, reduced in a reducing zone (900-1000°) with charcoal and gases from the carbonization of firewood or cane bagasse, pelletized by rehydration of the hemihydrate to the dihydrate, which is reduced to CaS (with CO2 and water recovery), and cooled, milled, and wetted to make a slurry. The liquid and solid phases are separated by centrifuges and pressurized filters, and the clear liquid containing soluble Ca(HS)2 is fed under pressure to a 2nd reactor receiving CO2 to cause formation of H2s and impurity-free CaCO3 precipitate The H2S produced in the 2nd reactor is fed under pressure to the 1st reactor, wherein S is produced by the Claus-Chance process.

ΙC C01F011-08

CC**49-9** (Industrial Inorganic Chemicals)

qypsum calcium carbonate sulfur prodn ST

13397-24-5P, preparation

RL: PREP (Preparation)

(calcium carbonate and sulfur manufacture from)

IT 471-34-1P, preparation 7704-34-9P, preparation

RL: PREP (Preparation)

(preparation of, from gypsum)

ΙT **471-34-1P**, preparation **7704-34-9P**, preparation

RL: PREP (Preparation)

(preparation of, from gypsum)

RN471-34-1 HCA

CN Carbonic acid calcium salt (1:1) (8CI, 9CI) (CA INDEX NAME)

HO-C-OH

● Ca

7704-34-9 HCA Sulfur (8CI, 9CI) (CA INDEX NAME)

S

L90 ANSWER 39 OF 54 HCA COPYRIGHT 2004 ACS on STN 100:90660 Desulfurization by three-stage combustion. (Hitachi Shipbuilding and Engineering Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 58190606 A2 19831107 Showa, 5 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1982-71978 19820428.

Flue gas is desulfurized by a 3-stage combustion system having AΒ desulfurizing agent recovery unit, Claus S recovery unit, and a secondary desulfurizing apparatus A fuel is fed to the 1st and 2nd combustors and then further combusted in a 3rd combustor with a secondary air supply. The flue gas is then treated in the secondary desulfurizing apparatus A desulfurizing agent (CaCO3, Na2CO3) and a portion of the effluent from the Claus unit are fed to the 2nd combustor. The spent desulfurizing agent is retrieved from the 2nd combustor to convert to carbonate which is recycled. The effluent from the carbonate converter is

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treated in a Claus unit to recover S.
 IC
      F23C011-00; B01D053-34; C01B017-04; F23C006-04
      59-4 (Air Pollution and Industrial Hygiene)
      Section cross-reference(s): 49
 IT
      Flue gases
         (desulfurization of, 3-stage combustion system for, desulfurizing agent
         and sulfur recovery in)
 IT
      497-19-8, uses and miscellaneous
      RL: USES (Uses)
         (for flue gas desulfurization, recycling of)
 IT
      7704-34-9P, preparation
      RL: PREP (Preparation)
         (recovery of, from flue gas desulfurization)
IT
      497-19-8, uses and miscellaneous
      RL: USES (Uses)
         (for flue gas desulfurization, recycling of)
RN
      497-19-8 HCA
CN
     Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)
HO-C-OH
 ●2 Na
IT
     7704-34-9P, preparation
     RL: PREP (Preparation)
         (recovery of, from flue gas desulfurization)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
L90 ANSWER 40 OF 54 HCA COPYRIGHT 2004 ACS on STN
98:18862 Recovery of calcium carbonate and sulfur from FGD scrubber
     waste. Arganbright, R. P.; Huang, P.; Benner, G. S.; Mandelik, B. G.;
     Roche, T. S. (Pullman Kellogg, Houston, TX, USA). Report,
     EPA-600/7-82-015; Order No. PB82-227729, 143 pp. Avail. NTIS From: Gov.
     Rep. Announce. Index (U. S.) 1982, 82(20), 4091 (English) 1982.
     The key process steps in the proprietary Kel-S process are demonstrated
     for recovering CaCO3 and S from lime/limestone flue gas desulfurization
     (FGD) scrubber waste. The steps are: reduction of the waste to CaS (using
     coal as the reducing agent), carbonation of the CaS to generate
     H2S and CaCO3, and recovery of precipitated CaCO3 from inerts
     (coal ash).
CC
     49-5 (Industrial Inorganic Chemicals)
     Section cross-reference(s): 59
     flue gas desulfurization; sulfur recovery; calcium
ST
     carbonate recovery
IT
     Flue gases
        (desulfurization of, calcium carbonate and sulfur
        recovery from)
     471-34-1P, preparation 7704-34-9P, preparation
ΙT
```

RL: PREP (Preparation)

(recovery of, from limestone flue gas desulfurization scrubber waste) 471-34-1P, preparation 7704-34-9P, preparation

IT 471-34-1P, preparation RL: PREP (Preparation)

(recovery of, from limestone flue gas desulfurization scrubber waste)

RN 471-34-1 HCA

CN Carbonic acid calcium salt (1:1) (8CI, 9CI) (CA INDEX NAME)

но- с- он О

Ca

RN 7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

L90 ANSWER 41 OF 54 HCA COPYRIGHT 2004 ACS on STN 97:8588 Removing hydrogen sulfide from gas streams. Gowdy, Hugh W.; Fenton, Donald M. (Union Oil Co., USA). U.S. US 4325936 A 19820420, 11 pp. Cont.-in-part of U.S. Ser. No. 50,193, abandoned. (English). CODEN: USXXAM. APPLICATION: US 1981-233888 19810212. PRIORITY: US 1979-50193 19790620.

AB A H2S removal and conversion method is described, in which a H2S-containing gas stream is contacted with a regenerable washing solution containing solubilized V thiocyanate ions, a carboxylate complexing agent, and one or more water-soluble quinones capable of solubilizing tetravalent V. The molar ratio of V to quinone(s) in the washing solution is selected to substantially reduce or eliminate the formation of contaminant sulfate salts. The absorbed H2S is converted to elemental S which, after oxidative regeneration of the washing solution, is separated from the regenerated solution

Thus, a gas containing H2S 1, CO2 7, and N2 92 mol% was washed for 16.8 days at 60°F and 1 atm with 0.037 L of washing solution/L of H2S-containing gas. The washing solution contained (in g/L): V 4.2, di-Na salt of 9,10-anthraquinone disulfonic acid (ADA, di-Na salt) 1.4, Na 1-hydroxybenzene-4-sulfonate (PSA) 22, NaSCN 87, Na citrate 13-16, Na carbonate 24, Na2S2O3 <0.2, and Na2SO4 1.3-2.6. The rates of change (g/L day) were: V 0.0, ADA di-Na salt 0.0, PSA Na salt 0.0, Na citrate loss 0.16, Na carbonate 0.0, Na2S2O3 0.0, Na2SO4 gain 0.07. The mol. ratios were V/ADA 24 and V/PSA 0.74. The amount of H2S converted was: to S2O32- salts 0.0 and SO42- salts 0.3%.

IC C01B017-05

NCL 423573000R

CC **49-10** (Industrial Inorganic Chemicals) Section cross-reference(s): 48

ST hydrogen sulfide removal gas; quinone solubilizer vanadium; sulfur recovery

TT 7704-34-9P, preparation

RL: PREP (Preparation)

(recovery of, from hydrogen sulfide-containing gases,

```
recyclable washing solution for)
IT
     994-36-5 1300-51-2 60553-45-9
     RL: USES (Uses)
        (solubilization by, of tetramine vanadium)
IT
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (recovery of, from hydrogen sulfide-containing gases,
        recyclable washing solution for)
     7704-34-9 HCA
RN
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
```

S

TT 994-36-5 RL: USES (Uses) (solubilization by, of tetramine vanadium) RN

1,2,3-Propanetricarboxylic acid, 2-hydroxy-, sodium salt (9CI) (CA INDEX CNNAME)

$$\begin{array}{c} {\rm CO_2H} \\ | \\ {\rm HO_2C-CH_2-C-CH_2-CO_2H} \\ | \\ {\rm OH} \end{array}$$

x Na

L90 ANSWER 42 OF 54 HCA COPYRIGHT 2004 ACS on STN 95:171827 Conversion of alkali metal sulfate to the carbonate. Sheth, Atul C. (United States Dept. of Energy, USA). U. S. Pat. Appl. US 80726 19800313, 18 pp. Avail. NTIS Order No. PAT-APPL-080 726. (English). CODEN: XAXXAV. APPLICATION: US 1979-80726 19791001.

A process is described for converting K2SO4 to K2CO3 in which a mixture of AΒ K2SO4 and CaO are reacted at $700-800^{\circ}$, with a gaseous mixture having a minor amount of H2 and/or CO in a diluent with the CaO being present in an amount <20 weight% of the K2SO4 to produce an aqueous mixture of K2S, KHS, KOH, and

CaS and a gaseous mixture of steam and H2S. The K and Ca salts are quenched to produce an aqueous slurry of soluble K salts and insol. Ca salts and a

mixture of steam and H2S. The Ca salts are separated from the solution The Ca salts are dried to produce CaS, Ca bisulfide, and steam and the CaS and Ca bisulfide are converted to the oxide and recycled. The soluble K salts are carbonated to produce K2CO3 which is concentrated and the precipitated crystals separated The S-containing compds. are treated further, e.g., by the Claus process.

CC 48-8 (Unit Operations and Processes) Section cross-reference(s): 49

potassium sulfate conversion carbonate; sulfur recovery MHD power plant

IT7778-80-5, reactions RL: RCT (Reactant); RACT (Reactant or reagent)

```
(conversion of, to carbonate)
 IT
      584-08-7P
      RL: PREP (Preparation)
         (manufacture of, from potassium sulfate for MHD power plant)
 IT
      7704-34-9P, preparation
      RL: PREP (Preparation)
         (recovery of, from coal-fired MHD power plant)
 IT
      584-08-7P
      RL: PREP (Preparation)
         (manufacture of, from potassium sulfate for MHD power plant)
      584-08-7 HCA
 RN
     Carbonic acid, dipotassium salt (8CI, 9CI) (CA INDEX NAME)
CN
HO-C-OH
 ●2 K
ΙT
     7704-34-9P, preparation
     RL: PREP (Preparation)
         (recovery of, from coal-fired MHD power plant)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
L90 ANSWER 43 OF 54 HCA COPYRIGHT 2004 ACS on STN
95:120132 Aqueous carbonate process design study. McKisson, R. L.;
     Bauerle, G. L.; Bodine, J. E.; Rennick, R. D.; Stewart, A. E.; Tsang, S.
     (Energy Syst. Group, Rockwell Int. Corp., Canoga Park, CA, USA). Report,
     EPRI-CS-1574, 466 pp. Avail. NTIS From: Energy Res. Abstr. 1981, 6(4),
     Abstr. No. 4483 (English) 1980.
AB
     The process involves a spray dryer in which the flue gas is scrubbed and
     its S removed as a dry Na2SO3/Na2SO4 plus Na2CO3 mixture This is followed
     by a regeneration process which involves the reduction of the SO32-/SO42- to
     S2- in a molten salt bed. This program involved studies of the
     molten salt reduction step and the conversion of an aqueous solution of the
     melt to a Na2CO3 scrubbing solution A process flow diagram and a material
     balance were developed based on coal-reduction process data. The reduction
     process was more difficult to operate with coal than with coke. The
     greater ash content of the coal is a major factor in determining the operating
     temperature of the reducer, with the result that the coal-reduction required a
bed
     temperature .apprx.55° higher than that for coke-reduction
CÇ
     59-2 (Air Pollution and Industrial Hygiene)
     Section cross-reference(s): 49, 60
     Flue gases
IT
        (desulfurization of, regeneration of waste from, reduction by coal in
        molten salt bed in)
_{\rm IT}
     Coal
     RL: OCCU (Occurrence)
        (flue gas desulfurization waste reduction by, in salt bed, for
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```
sulfur recovery and regeneration of scrubber solution)
 IT
      Waste solids
         (from flue gas desulfurization, reduction by coal in molten salt
         bath for sulfur recovery and regeneration of scrubber solution)
IT
      Salts, uses and miscellaneous
      RL: USES (Uses)
         (molten, coal in, flue gas desulfurization waste treatment
        by, for sulfur recovery and regeneration of scrubber solution)
IT
      497-19-8, uses and miscellaneous
     RL: USES (Uses)
         (flue gas desulfurization scrubber solution containing, regeneration of)
     7704-34-9P, preparation
     RL: IMF (Industrial manufacture); PREP (Preparation)
         (manufacture of, in reduction of sulfite-sulfate residue from
desulfurization
        scrubbing solution, coal in molten salt bath in)
IT
     7757-82-6, reactions
                             7757-83-7
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reduction of, from spent desulfurization scrubber solution, coal in
        molten salt bed in)
IT
     497-19-8, uses and miscellaneous
     RL: USES (Uses)
        (flue gas desulfurization scrubber solution containing, regeneration of)
RN
     497-19-8 HCA
     Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)
CN
HO-C-OH
 ●2 Na
     7704-34-9P, preparation
     RL: IMF (Industrial manufacture); PREP (Preparation)
        (manufacture of, in reduction of sulfite-sulfate residue from
desulfurization
        scrubbing solution, coal in molten salt bath in)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
```

L90 ANSWER 44 OF 54 HCA COPYRIGHT 2004 ACS on STN 94:5538 Recovering calcium and sulfur compounds from a metallurgical waste material. Collin, Per Harald (Swed.). PCT Int. Appl. WO 8000831 19800501, 12 pp. (English). CODEN: PIXXD2. APPLICATION: WO 1979-SE200 19791004.

AB Waste material containing CaO and CaS such as may be formed when S is removed from raw iron with lime or from preredn. of an iron ore with some grade of coal is treated. The CaO-CaS-containing waste is suspended in water for treatment with H2S until the partial pressure of H2S is 0.5-1.5 bar above the suspension and Ca(HS)2 is in solution After the insol. material is removed, the solution is treated with CO2 at a rate that assures that the

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gaseous product contains H2S with 10-30 volume% CO2 and CaCO3 is precipitated
 IC
      C01F011-10; C01F011-18
      49-9 (Industrial Inorganic Chemicals)
 CC
 ST
      iron ore preredn waste; hydrogen sulfide recovery;
      calcium carbonate recovery
 ΙT
      Wastes
         (from iron ore preredn., calcium and sulfur compound
         recovery from)
 IT
      Iron ores, reactions
      RL: RCT (Reactant); RACT (Reactant or reagent)
         (preredn. of, waste solids from, calcium and sulfur compound
        recovery from)
IT
      471-34-1P, preparation
     RL: PREP (Preparation)
         (preparation of, from iron ore preredn. waste solids)
IT
     7704-34-9P, preparation
                               7783-06-4P, preparation
     RL: PREP (Preparation)
        (recovery of, from iron ore preredn. waste solids)
IT
     471-34-1P, preparation
     RL: PREP (Preparation)
         (preparation of, from iron ore preredn. waste solids)
RN
     471-34-1 HCA
CN
     Carbonic acid calcium salt (1:1) (8CI, 9CI) (CA INDEX NAME)
HO-C-OH
  Ca
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (recovery of, from iron ore preredn. waste solids)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
L90 ANSWER 45 OF 54 HCA COPYRIGHT 2004 ACS on STN
93:243144 Rare anthropogenic and natural particles suspended in deep ocean
     waters. Jedwab, Jacques (Lab. Geochim., Univ. Libre Bruxelles, Brussels,
     B-1050, Belg.). Earth and Planetary Science Letters, 49(2), 551-64
     (English) 1980. CODEN: EPSLA2. ISSN: 0012-821X.
     Four hundred membrane filters from the Atlantic and Pacific deep waters
AΒ
     were scanned for colored and opaque particles by a method integrating the
     light and the scanning electron microscopes, the electron microprobe and
     the micro-x-ray diffraction camera. About 40 different types of particles
     were found. A tentative classification of the particles and of their
     morphol. and chemical varieties is presented, according to their
     anthropogenic or natural origins. The particles fall into 5 groups: (1)
     particle species exclusively produced by man's activities (brass, cobalt
     aluminate); (2) particles exclusively produced by natural processes (like
     amphibole, ilmenite); (3) particles rarely produced by natural processes,
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but massively so by man (like metallic Cu, Cr2O3); (4) a large group of particles massively produced by nature and by man (like magnetite spherules, malachite); (5) a very small group of particles whose origin could not be attributed at present. Very few entirely new compds. from the point of view of continental mineralogy were encountered. 53-5 (Mineralogical and Geological Chemistry)

IT Carbonates, occurrence Silicates, occurrence Sulfates, occurrence Sulfides, occurrence

RL: OCCU (Occurrence)

(suspended particles of, in deep waters, of Atlantic and Pacific oceans)

IT Carbonates, occurrence RL: OCCU (Occurrence)

> (hydrogen, suspended particles of, in deep waters, of Atlantic and Pacific oceans)

ΙT **7704-34-9P**, occurrence

RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

IT 1309-38-2, occurrence 1310-14-1 1317-60-8, occurrence 1317-80-2 7429-90-5, occurrence 7439-89-6, 1317-92-6 **1319-53-5** 7440-50-8, occurrence 12597-68-1, occurrence 12597-71-6, occurrence 15860-78-3 RL: OCCU (Occurrence)

(suspended particles of, in deep waters, of Atlantic and Pacific oceans)

IT 7704-34-9P, occurrence

RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

RN 7704-34-9 HCA

Sulfur (8CI, 9CI) (CA INDEX NAME) CN

S

IT 1319-53-5

RL: OCCU (Occurrence)

(suspended particles of, in deep waters, of Atlantic and Pacific oceans)

1319-53-5 HCA RN

CN Malachite (Cu2(CO3)(OH)2) (9CI) (CA INDEX NAME)

HO-Cu-O-C-O-Cu-OH

L90 ANSWER 46 OF 54 HCA COPYRIGHT 2004 ACS on STN 89:45792 Treatment of an alkali metal sulfide solution. Rennick, Robert D. (Rockwell International Corp., USA). Ger. Offen. DE 2741613 19780316, 24 pp. (German). CODEN: GWXXBX. APPLICATION: DE 1977-2741613 19770915.

The solution is treated to recovery S. An alkaline solution with >60 mol % of AΒ

alkali metal sulfide is contacted in a neutralization zone with a H2S-containing gas to form a solution of reduced alkalinity containing a large amount of an

John Calve EIC- 1700

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alkali metal bisulfide and a lesser amount of the carbonate or
      bicarbonates. The solution is then contacted with a gas containing a low
 concentration
      of CO2 to yield carbonate crystals free of sulfides and sulfites
      and a product gas high in H2S, 35-65% of which is used in the
      neutralization zone and the rest treated in a Claus facility to produce S.
      The carbonate product stream is decomposed thermally to yield a
      waste gas high in CO2 and an aqueous slurry containing mainly carbonates
      . The CO2 is used in the carbonating step.
 IC
      C01B017-06
 CC
      49-1 (Industrial Inorganic Chemicals)
      sulfide alkali metal decompn; sulfur recovery
 ST
 IT
      124-38-9, reactions
      RL: RCT (Reactant); RACT (Reactant or reagent)
         (reaction of, with alkali metal sulfide, in sulfur
         recovery)
 ΙT
      7704-34-9P, preparation
      RL: PREP (Preparation)
         (recovery of, from alkali metal sulfide solns., carbon dioxide in)
ΙT
      7783-06-4D, alkali metal salts
     RL: USES (Uses)
         (sulfur recovery from, carbon dioxide in)
IT
     124-38-9, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
         (reaction of, with alkali metal sulfide, in sulfur
RN
     124-38-9 HCA
     Carbon dioxide (8CI, 9CI) (CA INDEX NAME)
CN
o = c = o
     7704-34-9P, preparation
IT
     RL: PREP (Preparation)
        (recovery of, from alkali metal sulfide solns., carbon dioxide in)
RN
     7704-34-9 HCA
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
L90 ANSWER 47 OF 54 HCA COPYRIGHT 2004 ACS on STN
87:41450 Removal of hydrogen sulfide from natural gas to obtain elemental
     sulfur. Kalymon, Ya. A.; Mel'nik, V. F.; Konovalenko, Z. L.; Pogonets, O.
     A.; Tomin, I. I. (USSR). Tezisy Dokl. Vses. Nauchno-Tekh. Konf. Tekhnol.
     Neorg. Veshchestv Miner. Udobr., 9th, Volume 1, 210-11. Editor(s):
     Amirova, S. A. Permsk. Politekh. Inst.: Perm, USSR. (Russian)
     1974. CODEN: 34PSAU.
     Optimum condition were discussed for removal of H2S from natural gas by
AΒ
     chemisorption in a Na2CO3 solution and preparation of elemental S by oxidn
     . of H2S with organic catalysts dissolved in the Na2CO3 solution
     51-4 (Fossil Fuels, Derivatives, and Related Products)
CC
     Section cross-reference(s): 49, 67
     natural gas hydrogen sulfide removal; sulfur prepn hydrogen
ST
     sulfide oxidn; catalyst hydrogen sulfide
     oxidn
IT
     Chemisorption
```

Hertzog

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(of hydrogen sulfide in sodium carbonate, for sweetening of
        natural gas)
IT
     Oxidation catalysts
         (soluble organic, for recovery of sulfur from sweetening
        of natural gas)
IT
      497-19-8, uses and miscellaneous
     RL: USES (Uses)
        (in removal of hydrogen sulfide from natural gas and associated
        sulfur recovery)
IT
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (preparation of, removal of hydrogen sulfide from natural gas in relation
IT
     497-19-8, uses and miscellaneous
     RL: USES (Uses)
        (in removal of hydrogen sulfide from natural gas and associated
        sulfur recovery)
     497-19-8 HCA
     Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)
HO-C-OH
 ●2 Na
IT
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (preparation of, removal of hydrogen sulfide from natural gas in relation
        to)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
L90 ANSWER 48 OF 54 HCA COPYRIGHT 2004 ACS on STN
86:142270 Study of reagent conditions for autoclave melting of sulfur. Tsap,
     S. M.; Mel'nik, V. F. (USSR). Tezisy Dokl. Vses. Nauchno-Tekh. Konf.
     Tekhnol. Neorg. Veshchestv Miner. Udobr., 9th, Volume 1, 203-4.
     Editor(s): Amirova, S. A. Permsk. Politekh. Inst.: Perm, USSR. (Russian)
     1974. CODEN: 34PSAU.
    To optimize the melting of S, the relation between reagent conditions and
AΒ
     composition and the properties of the concs. was investigated. The optimum
     consumption of Na2CO3 corresponds to the amount necessary for binding Ca2+
     and Mg2+ contained in the water added during melting. Na5P3O10
     consumption depends on the overall surface area of the barren rock. The
     efficiency of extracting S is decreased by sludge formation.
     49-1 (Industrial Inorganic Chemicals)
CC
    sulfur melting autoclave; sodium carbonate sulfur melting;
ST
    triphosphate sulfur ore extn
IT
     497-19-8, uses and miscellaneous 7758-29-4
    RL: USES (Uses)
        (in sulfur recovery from ore concs., by melting)
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7704-34-9P, preparation
      RL: PREP (Preparation)
         (recovery of, from ore concs. by melting, optimization of)
 IT
      497-19-8, uses and miscellaneous
      RL: USES (Uses)
         (in sulfur recovery from ore concs., by melting)
 RN
      497-19-8 HCA
      Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)
 CN
    O
 HO-C-OH
 ●2 Na
 ΙT
     7704-34-9P, preparation
     RL: PREP (Preparation)
         (recovery of, from ore concs. by melting, optimization of)
     7704-34-9 HCA
RN
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
L90 ANSWER 49 OF 54 HCA COPYRIGHT 2004 ACS on STN
83:47666 Recovery of sulfur dioxide. Shah, Indravadan S.
     (Chemical Construction Corp., USA). Can. CA 949289 19740618, 13
     pp. (English). CODEN: CAXXA4. APPLICATION: CA 1971-114866 19710604.
     Waste gases containing SO2 are scrubbed with a Na sulfite solution to form a Na
AΒ
     bisulfite solution, part of which is reacted with Ca and/or MgCO3 to form a
     precipitate which is heated, forming a recoverable S-containing product and
solid
     particles of Mg and/or CaO. The remaining part of the bisulfite solution is
     regenerated with Na2CO3 and recycled for further waste gas scrubbing.
     Thus, flue gas containing SO2 2000 ppm was delivered at 3010 m3/min and
     scrubbed to remove .apprx.94% of the initial SO2 content. After complete
     processing S was recovered at a rate of 5.45 kg/min.
     59-2 (Air Pollution and Industrial Hygiene)
     Section cross-reference(s): 49
ST
     sulfur recovery waste gas
IT
     Flue gases
        (sulfur dioxide removal from, by sodium bisulfite scrubber solution and
        magnesium carbonate, sulfur recovery in)
IT
     546-93-0
     RL: OCCU (Occurrence)
        (desulfurization scrubber solution treatment by, sulfur
        recovery in)
IT
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (recovery of, in sulfur dioxide removal from flue
        gas by sodium bisulfite scrubber solution and magnesium carbonate
     7446-09-5, uses and miscellaneous
IT
     RL: REM (Removal or disposal); PROC (Process)
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```
(removal of, from flue gas by sodium bisulfite scrubber solution and
          magnesium carbonate, sulfur recovery in)
      7631-90-5
 IT
      RL: OCCU (Occurrence)
          (sulfur dioxide removal by scrubber solution containing, from flue gas,
          sulfur recovery in)
 IT
      546-93-0
      RL: OCCU (Occurrence)
          (desulfurization scrubber solution treatment by, sulfur
         recovery in)
 RN
      546-93-0 HCA
 CN
      Carbonic acid, magnesium salt (1:1) (8CI, 9CI) (CA INDEX NAME)
 HO-C-OH
 IT
      7704-34-9P, preparation
      RL: PREP (Preparation)
         (recovery of, in sulfur dioxide removal from flue
         gas by sodium bisulfite scrubber solution and magnesium carbonate
      7704-34-9 HCA
RN
CN
      Sulfur (8CI, 9CI) (CA INDEX NAME)
S
L90 ANSWER 50 OF 54 HCA COPYRIGHT 2004 ACS on STN
80:99731 Chemistry of the molten carbonate process for
     sulfur oxides removal stack gases. Yosim, S. J.; Grantham, L. F.;
     McKenzie, D. E.; Stegmann, G. C. (At. Int. Div., Rockwell Int. Corp.,
     Canoga Park, CA, USA). Advances in Chemistry Series, 127, 174-82
     (English) 1973. CODEN: ADCSAJ. ISSN: 0065-2393. In the molten carbonate process, a molten eutectic mixture of Li,
AB
     Na, and K carbonates removes S oxides from power plant stack
     gases. The resulting molten solution of alkali metal sulfites, sulfates, and
     unreacted carbonate is regenerated in a 2-step process to the
     alkali carbonate for recycling. H2S, which is evolved in the
     regeneration step, is converted to S in a conventional Claus plant. A
     10-MW pilot plant of the process has been constructed at the Consolidated
     Edison Arthur Kill Station on Staten Island, and startup is underway.
     59-2 (Air Pollution and Industrial Hygiene)
     carbonate sulfur oxide removal; stack gas sulfur oxide removal
ST
     Carbonates, uses and miscellaneous
     RL: USES (Uses)
        (sulfur oxide removal by molten, from flue gas)
IΤ
     Flue gases
        (sulfur oxide removal from, molten
        carbonates in)
ΙT
     7704-34-9P, preparation
     RL: PREP (Preparation)
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(recovery of, in sulfur oxide removal from flue gas by molten
        carbonates)
     7446-09-5, uses and miscellaneous 7446-11-9, uses and miscellaneous
     RL: REM (Removal or disposal); PROC (Process)
        (removal of, from flue gas, molten carbonates in)
     497-19-8, uses and miscellaneous 554-13-2
     584-08-7
     RL: USES (Uses)
        (sulfur oxide removal by molten, from flue gas)
IT
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (recovery of, in sulfur oxide removal from flue gas by molten
        carbonates)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
     497-19-8, uses and miscellaneous 554-13-2
     584-08-7
     RL: USES (Uses)
        (sulfur oxide removal by molten, from flue gas)
     497-19-8 HCA
RN
     Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)
CN
HO-C-OH
 ●2 Na
RN
     554-13-2 HCA
CN Carbonic acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)
   0
HO-C-OH
•2 Li
RN 584-08-7 HCA
CN Carbonic acid, dipotassium salt (8CI, 9CI) (CA INDEX NAME)
   0
HO-C-OH
```

●2 K

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L90 ANSWER 51 OF 54 HCA COPYRIGHT 2004 ACS on STN
80:52188 Treating liquid waste containing sulfur.
     Ninagawa, Hiroshi Jpn. Kokai Tokkyo Koho JP 48063967 19730905
     Showa, 4 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1971-99728
     19711209.
     The spent alkaline waste solution used to remove H2S was recovered. CO2 was
AB
     bubbled into the solution at a temperature above the decomposition temperature
of alkali H
     carbonate to react with alkali H sulfide or alkali sulfide. The
     H2S evolved with CO2 was removed from the solution and was absorbed in a
     solution for recovery of elementary S.
NCL
    13(7)A21
CC
     60-2 (Sewage and Wastes)
IT
     124-38-9, uses and miscellaneous
     RL: USES (Uses)
        (hydrogen sulfide removal by, from desulfurizer scrubber solution, sulfur
        recovery in)
IT
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (recovery of, from desulfurizer scrubber solution, carbon dioxide in)
IΤ
     124-38-9, uses and miscellaneous
     RL: USES (Uses)
        (hydrogen sulfide removal by, from desulfurizer scrubber solution, sulfur
        recovery in)
RN
     124-38-9 HCA
CN
     Carbon dioxide (8CI, 9CI) (CA INDEX NAME)
0 = c = 0
IT
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (recovery of, from desulfurizer scrubber solution, carbon dioxide in)
ВN
     7704-34-9 HCA
CN
    Sulfur (8CI, 9CI) (CA INDEX NAME)
S
L90 ANSWER 52 OF 54 HCA COPYRIGHT 2004 ACS on STN
79:106507 Sulfur from Claus-process off-gases. Peter, Siegfried
    Ger. Offen. DE 2165646 19730705, 14 pp. (German). CODEN:
    GWXXBX. APPLICATION: DE 1971-2165646 19711230.
    H2S and SO2 from the off-gases of the S separator of the 2nd
    catalyst step of the Claus-process were converted to S by washing at
    120° with polyalkylene glycols or their ethers, e.g. triethylene
    glycol (I) or hexaethylene glycol monobutyl ether in the presence of
    alkanolamines, e.g. (HOCH2CH2)3N or [MeCH(OH)CH2]2NH (II),
    triethylenetetramine, or KHCO3 as catalyst. Thus, a gas containing N 70,
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IC C01B

CC 49-1 (Industrial Inorganic Chemicals)

of clear yellow color after solidification.

solution containing 8% II in I to give a gas containing 0.5 volume % H2S and

H2O(g) 25, H2S 3.5, and SO2 1.5 volume $\frac{1}{8}$ was passed through a

traces SO2. The liquid S formed was drawn off the bottom and was

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Section cross-reference(s): 59
      sulfur Claus process gas; hydrogen sulfide
 ST
      sulfur; oxide sulfur hydrogen sulfide;
      polyalkylene glycol gas washing; ether polyalkylene glycol washing;
      alkanolamine catalyst sulfur; ethyleneamine catalyst
      sulfur; amine catalyst sulfur; potassium
      carbonate catalyst sulfur
 IT
      Catalysts and Catalysis
         (amines and potassium carbonate, for hydrogen sulfide
         -sulfur dioxide reaction)
 IT
      102-71-6, uses and miscellaneous
                                         110-97-4 112-24-3 141-43-5, uses
      and miscellaneous 298-14-6 622-40-2
      RL: CAT (Catalyst use); USES (Uses)
         (catalysts, in sulfur manufacture from Claus process gases)
 IT
      7704-34-9P, preparation
      RL: PREP (Preparation)
         (from hydrogen sulfide- and sulfur dioxide-containing
        gases, by washing with polyalkylene glycols containing alkaline catalysts)
 TT
     112-27-6 112-60-7
                          1072-40-8 1191-91-9 4403-55-8 4792-15-8
      25322-68-3 25322-69-4
      RL: USES (Uses)
         (in sulfur manufacture from Claus-process gases, washing with)
IT
     7446-09-5, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
         (with hydrogen sulfide, for sulfur manufacture from
        Claus process gases)
ΙT
     7783-06-4, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (with sulfur dioxide, for sulfur manufacture from Claus
        process gases)
IT
     298-14-6
     RL: CAT (Catalyst use); USES (Uses)
        (catalysts, in sulfur manufacture from Claus process gases)
RN
     298-14-6 HCA
     Carbonic acid, monopotassium salt (8CI, 9CI) (CA INDEX NAME)
HO-C-OH
ΙT
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (from hydrogen sulfide- and sulfur dioxide-containing
        gases, by washing with polyalkylene glycols containing alkaline catalysts)
     7704-34-9 HCA
RN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
CN
L90 ANSWER 53 OF 54 HCA COPYRIGHT 2004 ACS on STN
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74:143938 Converting soda slag from a blast furnace to useful products.

S

Hertzog

(Kloeckner-Humboldt-Deutz A.-G.). Ger. Offen. DE 1934308 19710211, 14 pp. (German). CODEN: GWXXBX. APPLICATION: DE 1969-1934308 19690707.

- AB A continuous flow process is described whereby obnoxious and poisonous H2O-soluble and easily decomposable constituents of a slag can be removed and converted to other products. Specifically, slag is ground with H2O. The solution which contains primarily Na2S is separated from the insol. portion and is treated with CO2 at elevated temps. at a high pressure. H2S is liberated from the solution and NaHCO3 is formed in solution. The H2S is oxidized to elemental S and NaHCO3 isolated from the solution and converted to Na2CO3 by calcination.
- IC C01D; C01B; C21C
- CC 49 (Industrial Inorganic Chemicals)
- ST blast furnace sulfide slag conversion; sulfur recovery; sodium carbonate recycling
- IT Slags

(sodium sulfide removal from blast-furnace, for sodium

carbonate and sulfur recovery)

IT 497-19-8P, preparation 7704-34-9P, preparation

RL: PREP (Preparation)

(recovery of, from sodium sulfide from blast-furnace slags)

IT 1313-82-2

RL: REM (Removal or disposal); PROC (Process)

(removal of, from blast-furnace slags for recovery of sodium carbonate and sulfur)

IT 497-19-8P, preparation 7704-34-9P, preparation

RL: PREP (Preparation)

(recovery of, from sodium sulfide from blast-furnace slags)

RN 497-19-8 HCA

CN Carbonic acid disodium salt (8CI, 9CI) (CA INDEX NAME)

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•2 Na

RN 7704-34-9 HCA

CN Sulfur (8CI, 9CI) (CA INDEX NAME)

S

L90 ANSWER 54 OF 54 HCA COPYRIGHT 2004 ACS on STN 74:55749 Carbon oxide regenerant for sulfur recovery from molten salts. Grantham, Le Roy F. (North American Rockwell Corp.). U.S. US 3551108 19701229, 8 pp. Division of U.S. 3,438,734 (English). CODEN: USXXAM. APPLICATION: US 1968-779175 19681126.

AB A method is described for direct recovery of elemental S from the molten salt mixture used for desulfurizing flue gases according to the process of U.S. 3,438,734. The molten salt mixture containing

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alkali metal sulfites is treated at 400-50^{\circ} with H and (or) CO to
      reduce partially the sulfites to sulfides. The melt is then
      treated with CO2 whereby elemental S vapor and a melt of alkali metal
     carbonates are formed. The molten carbonates
     are recycled to the flue gas desulfurization step and the S is condensed
      from the vapor as product. The method avoids formation of H2S
      as an intermediate requiring conversion to S.
     C01D; C01B
IC
NCL 023224000
CC
     49 (Industrial Inorganic Chemicals)
ST
     sulfur recovery; flue gases desulfurizing; gases flue
     desulfurizing; desulfurizing flue gases
IT
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (recovery of, from alkali metal sulfites by reduction with carbon oxides
        and hydrogen)
IT
     124-38-9, reactions
                          630-08-0, reactions
                                                 1333-74-0, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reduction by, of alkali metal sulfites for sulfur recovery)
IT
     7704-34-9P, preparation
     RL: PREP (Preparation)
        (recovery of, from alkali metal sulfites by reduction with carbon oxides
        and hydrogen)
RN
     7704-34-9 HCA
CN
     Sulfur (8CI, 9CI) (CA INDEX NAME)
S
ΤТ
     124-38-9, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reduction by, of alkali metal sulfites for sulfur recovery)
RN
     124-38-9 . HCA
CN
     Carbon dioxide (8CI, 9CI) (CA INDEX NAME)
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